WEST Search History

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DATE: Thursday, October 14, 2004

Hide?	<u>Set Nan</u>	<u>ne Query</u>	Hit Count
	DB=PC	GPB, USPT, USOC, EPAB, JPAB, DWPI; PLUR = YES;	OP=ADJ
П	L23	(L19 AND olfactory)	62
	L22	(L19 AND inhale)	7
	L21	L19 AND nasal administration	23
	L20	L19 AND inhalation therapy	8
	L19	L18 AND coronary artery disease	805
	L18	FGF-1 OR FGF-2 OR aFGF OR bFGF OR VEGF	12449
	L17	L16 NOT Ashkenazi-Avi-J.IN.	173
WV 25	L16	L15 NOT Ashkenazi-Avi.IN.	173
	L15	L14 NOT Rosen-Craig.IN.	245
	L14	L13 NOT Rosen-Craig-A.IN.	245
	L13	L12 AND nasal	281
	L12	L11 AND inhalation	345
	L11	(L7 AND coronary artery disease)	540
	L10	L9 AND coronary artery disease	26
	L9	L8 AND inhalation	532
	L8	L6 AND growth factor	2460
	L7	530/300,350,399.CCLS.	18515
	L6	514/2.CCLS.	6335
	L5	Franco.IN.	6556
	L4	Franco-W.IN.	4
	L3	Franco-W-P.IN.	3
	L2	Franco-Wayne.IN.	0
	L1	(Franco-Wayne-P.IN.)	7

END OF SEARCH HISTORY

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Search Results - Record(s) 1 through 6 of 6 returned.

☐ 1. Document ID: US 20040167070 A1

Using default format because multiple data bases are involved.

L24: Entry 1 of 6

File: PGPB

Aug 26, 2004

PGPUB-DOCUMENT-NUMBER: 20040167070

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040167070 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor

and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: August 26, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

□ 2. Document ID: US 20040116349 A1

L24: Entry 2 of 6

File: PGPB

Jun 17, 2004

PGPUB-DOCUMENT-NUMBER: 20040116349

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040116349 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: June 17, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising

 $http://westbrs:9000/bin/gate.exe?f=TOC\&state=3pgbkl.25\&ref=24\&dbname=PGPB, USPT, U... \\ 10/14/04$

the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw, Desc

☐ 3. Document ID: US 20040023863 A1

L24: Entry 3 of 6

File: PGPB

Feb 5, 2004

PGPUB-DOCUMENT-NUMBER: 20040023863

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040023863 A1

TITLE: Methods of use growth factors for treating heart disease

PUBLICATION-DATE: February 5, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of <u>CPK-MB</u> levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMAC	Drawn Des

☐ 4. Document ID: US 20020058612 A1

L24: Entry 4 of 6

File: PGPB

May 16, 2002

PGPUB-DOCUMENT-NUMBER: 20020058612

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020058612 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: May 16, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

http://westbrs:9000/bin/gate.exe?f=TOC&state=3pgbkl.25&ref=24&dbname=PGPB,USPT,U... 10/14/04

US-CL-CURRENT: 514/2; 424/43

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of <u>CPK-MB</u> levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full Title Citation Front Review Classification D	ate Reference	Sequences	Attachments	Claims	10000	Draw, Des
☐ 5. Document ID: US 6759386 B2						

US-PAT-NO: 6759386

DOCUMENT-IDENTIFIER: US 6759386 B2

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

DATE-ISSUED: July 6, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Franco; Wayne P.

Rocky Hill

CT

06067

US-CL-CURRENT: 514/2; 514/12, 514/14, 514/8, 530/300

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

24 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

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Full	Title	Citation	Front	Review	Classification	Date	Reference	Claima	1-204C	Draw, Desi
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☐ 6. Document ID: US 20040167070 A1, WO 200177328 A1, AU 200155237 A, US 20020058612 A1, US 20040023863 A1, US 20040116349 A1, US 6759386 B2

L24: Entry 6 of 6

File: DWPI

Aug 26, 2004

DERWENT-ACC-NO: 2002-049148

DERWENT-WEEK: 200457

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TITLE: Treatment of heart disease brought on by e.g. myocardial infarction, unstable angina, thrombolytic therapy, bypass surgery or angioplasty, comprises multi-tiered administration of growth factors

INVENTOR: FRANCO, W P

PRIORITY-DATA: 2000US-195624P (April 6, 2000), 2001US-0828330 (April 6, 2001), 2003US-0730831 (December 9, 2003), 2003US-0731197 (December 9, 2003)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
US 20040167070 A1	August 26, 2004		000	A61K038/18
WO 200177328 A1	October 18, 2001	E	089	C12N015/12
AU 200155237 A	October 23, 2001		000	·
US 20020058612 A1	May 16, 2002		000	A61L009/04
US 20040023863 A1	February 5, 2004		000	A61K038/18
US 20040116349 A1	June 17, 2004		000	A61K038/18
US 6759386 B2	July 6, 2004		000	A61K038/18

INT-CL (IPC): $\underline{A61}$ \underline{K} $\underline{38/18}$; $\underline{A61}$ \underline{L} $\underline{9/04}$; $\underline{C07}$ \underline{K} $\underline{5/00}$; $\underline{C07}$ \underline{K} $\underline{14/52}$; $\underline{C12}$ \underline{N} $\underline{15/12}$

ABSTRACTED-PUB-NO: US20020058612A BASIC-ABSTRACT:

 ${\tt NOVELTY-Treatment\ of\ heart\ disease\ comprising\ multi-tiered\ administration\ of\ growth\ factors.}$

DETAILED DESCRIPTION - Systematic multi-tiered treatment of heart disease comprises delivery of therapeutic growth factor proteins (GFP) by:

- (a) oral inhalation of at least one dose of an effective amount of a first therapeutic GFP formulation in a patient displaying symptoms of heart disease;
- (b) monitoring levels of CPK-MB in the patient;
- (c) determining whether administration of the GFP formulation was effective in treating the symptoms;
- (d) administering one or ore additional doses of a second GFP formulation by a delivery method more invasive than oral inhalation; and
- (e) repeating steps (b)-(d) until there is a clinical indication of amelioration of the symptoms of heart disease in the patient, or until there is a contraindication to continued treatment.

INDEPENDENT CLAIMS are also included for the following:

- (1) administration of therapeutic amounts of GFP formulation for treatment of heart disease by inhalation;
- (2) monitoring clinical effectiveness of administration of a GFP formulation in the treatment of heart disease comprising:
- (i) performing an assay on a sample of biological fluid from a patient displaying symptoms of heart disease to determine the amount of $\underline{\text{CPK-MB}}$ present in the fluid;
- (ii) administering a therapeutic amount of GFP formulation to the patient; and

(iii) repeating steps (i) and (ii) until the assayed amount of $\underline{\text{CPK-MB}}$ in the biological fluid has decreased by an amount sufficient to indicate the clinical effectiveness of the administration of the GFP formulation.

ACTIVITY - Cardiant. Pellets containing 10 or 100 micro g basic fibroblast growth factor (bFGF) or placebo were placed on the epicardial surface in patients with a viable and ischemic myocardial area that could not be revascularized, during coronary artery bypass surgery. After 16 months patients were angina free with the exception of 3 people in the placebo group and 1 patient who received the 10 micro g pellet.

MECHANISM OF ACTION - Angiogenesis stimulator.

USE - The method is used for the treatment of heart disease, where the symptoms are chronic or acute, especially where the symptoms are brought on by myocardial infarction, unstable angina, acute anginal attack or reperfusion injury, preferably induced by thrombolytic therapy, bypass surgery or angioplasty (claimed).

ADVANTAGE - Inhalation is the least invasive method of delivering the growth factors to the lungs. Prior are invasive approaches have not been successful in promoting angiogenesis. The pericardial space serves as a drug delivery reservoir for delivery of therapeutic agents to the heart. Use of a catheter avoids the need for open chest surgery. Intravenous infusions are practical, low cost and can be used in a broad group of patients. Treatment can be repeated easily and may not require any special facilities.

ABSTRACTED-PUB-NO:

WO 200177328A EQUIVALENT-ABSTRACTS:

NOVELTY - Treatment of heart disease comprising multi-tiered administration of growth factors.

DETAILED DESCRIPTION - Systematic multi-tiered treatment of heart disease comprises delivery of therapeutic growth factor proteins (GFP) by:

- (a) oral inhalation of at least one dose of an effective amount of a first therapeutic GFP formulation in a patient displaying symptoms of heart disease;
- (b) monitoring levels of $\underline{CPK-MB}$ in the patient;
- (c) determining whether administration of the GFP formulation was effective in treating the symptoms;
- (d) administering one or ore additional doses of a second GFP formulation by a delivery method more invasive than oral inhalation; and
- (e) repeating steps (b)-(d) until there is a clinical indication of amelioration of the symptoms of heart disease in the patient, or until there is a contraindication to continued treatment.

INDEPENDENT CLAIMS are also included for the following:

- (1) administration of therapeutic amounts of GFP formulation for treatment of heart disease by inhalation;
- (2) monitoring clinical effectiveness of administration of a GFP formulation in the treatment of heart disease comprising:
- (i) performing an assay on a sample of biological fluid from a patient displaying symptoms of heart disease to determine the amount of CPK-MB present in the fluid;
- (ii) administering a therapeutic amount of GFP formulation to the patient; and

(iii) repeating steps (i) and (ii) until the assayed amount of $\underline{CPK-MB}$ in the biological fluid has decreased by an amount sufficient to indicate the clinical effectiveness of the administration of the GFP formulation.

ACTIVITY - Cardiant. Pellets containing 10 or 100 micro g basic fibroblast growth factor (bFGF) or placebo were placed on the epicardial surface in patients with a viable and ischemic myocardial area that could not be revascularized, during coronary artery bypass surgery. After 16 months patients were angina free with the exception of 3 people in the placebo group and 1 patient who received the 10 micro g pellet.

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ADVANTAGE - Inhalation is the least invasive method of delivering the growth factors to the lungs. Prior are invasive approaches have not been successful in promoting angiogenesis. The pericardial space serves as a drug delivery reservoir for delivery of therapeutic agents to the heart. Use of a catheter avoids the need for open chest surgery. Intravenous infusions are practical, low cost and can be used in a broad group of patients. Treatment can be repeated easily and may not require any special facilities.

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Search Results - Record(s) 1 through 7 of 7 returned.

☐ 1. Document ID: US 20040167070 A1

Using default format because multiple data bases are involved.

L1: Entry 1 of 7

File: PGPB

Aug 26, 2004

PGPUB-DOCUMENT-NUMBER: 20040167070

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040167070 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: August 26, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

☐ 2. Document ID: US 20040116349 A1

L1: Entry 2 of 7

File: PGPB

Jun 17, 2004

PGPUB-DOCUMENT-NUMBER: 20040116349

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040116349 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: June 17, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising

the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KMC | Draw Desi

☐ 3. Document ID: US 20040023863 A1

L1: Entry 3 of 7

File: PGPB

Feb 5, 2004

PGPUB-DOCUMENT-NUMBER: 20040023863

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040023863 A1

TITLE: Methods of use growth factors for treating heart disease

PUBLICATION-DATE: February 5, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

Franco, Wayne P.

Rocky Hill

CT

US

RULE-47

US-CL-CURRENT: 514/12

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims RMC Draw Des

☐ 4. Document ID: US 20020058612 A1

L1: Entry 4 of 7

File: PGPB

May 16, 2002

PGPUB-DOCUMENT-NUMBER: 20020058612

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020058612 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: May 16, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

US

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US-CL-CURRENT: 514/2; 424/43

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	FMMC	Draw Des
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L1: Entry 5 of 7

File: USPT

Jul 6, 2004

US-PAT-NO: 6759386

DOCUMENT-IDENTIFIER: US 6759386 B2

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

DATE-ISSUED: July 6, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Franco; Wayne P.

Rocky Hill

CT

06067

US-CL-CURRENT: 514/2; 514/12, 514/14, 514/8, 530/300

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

24 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title Citation	n Front	Review	Classification	Date	Reference		Claims	KWIC	Draw, De.
☐ 6. Docum	nent ID:	US 437	78347 A				***************************************	***************************************	***************************************

US-PAT-NO: 4378347

DOCUMENT-IDENTIFIER: US 4378347 A

TITLE: Composition for treating the heart for myocardial infarction

DATE-ISSUED: March 29, 1983

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Franco; Wayne P.

Wethersfield

CT

06109

US-CL-CURRENT: 424/565; 514/21, 514/777

ABSTRACT:

An effective dose of FGF for treatment of the heart is suspended in a slow release carrier and used in treatment of ischemic heart disease.

2 Claims, 0 Drawing figures Exemplary Claim Number: 1,2

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	K004C	Draw, De:
П	7 1	Docume	nt ID:	115 42	06100 4	***************************************	***************************************	***************************************	······································	······································	***************************************	······
		Docume:		US 42	96100 A	***************************************	***************************************		······································		······································	

US-PAT-NO: 4296100

DOCUMENT-IDENTIFIER: US 4296100 A

** See image for Certificate of Correction **

TITLE: Method of treating the heart for myocardial infarction

DATE-ISSUED: October 20, 1981

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Franco; Wayne P.

Wethersfield

CT

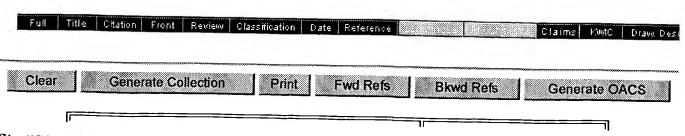
06109

US-CL-CURRENT: 424/565; 514/21

ABSTRACT:

The heart is treated with fibroblast growth factor to alleviate conditions caused by myocardial infarctions as by reducing the size of damaged heart areas. An effective dose of fibroblast growth factor when applied to the heart is found to increase blood flow in affected areas for a period of at least 4 hours and often more.

13 Claims, 0 Drawing figures Exemplary Claim Number: 1



Terms	Documents
(Franco-Wayne-P.IN.)	7

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Search Results - Record(s) 1 through 3 of 3 returned.

Document ID: US 20040167070 A1, WO 200177328 A1, AU 200155237 A, US 20020058612 A1, US 20040023863 A1, US 20040116349 A1, US 6759386 B2

Using default format because multiple data bases are involved.

L2: Entry 1 of 3

File: DWPI

Aug 26, 2004

DERWENT-ACC-NO: 2002-049148

DERWENT-WEEK: 200457

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: Treatment of heart disease brought on by e.g. myocardial infarction, unstable angina, thrombolytic therapy, bypass surgery or angioplasty, comprises multi-tiered

administration of growth factors

INVENTOR: FRANCO, W P

PRIORITY-DATA: 2000US-195624P (April 6, 2000), 2001US-0828330 (April 6, 2001),

2003US-0730831 (December 9, 2003), 2003US-0731197 (December 9, 2003)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
US 20040167070 A1	August 26, 2004		000	A61K038/18
WO 200177328 A1	October 18, 2001	E	089	C12N015/12
AU 200155237 A	October 23, 2001		000	,
US 20020058612 A1	May 16, 2002		000	A61L009/04
US 20040023863 A1	February 5, 2004		000	A61K038/18
US 20040116349 A1	June 17, 2004		000	A61K038/18
US 6759386 B2	July 6, 2004		000	A61K038/18

INT-CL (IPC): A61 K 38/18; A61 L 9/04; C07 K 5/00; C07 K 14/52; C12 N 15/12

Full	Title	Citation	Front	Review	Classification	Date	Reference		Claims	KWIC	Draw, Des

☐ 2. Document ID: US 4378347 A.

L2: Entry 2 of 3

File: DWPI

Mar 29, 1983

DERWENT-ACC-NO: 1983-36768K

DERWENT-WEEK: 198315

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TITLE: Fibroblast growth factor in slow-release carrier - for treating ischaemic

heart disease

INVENTOR: FRANCO, W P

PRIORITY-DATA: 1981US-0274722 (June 18, 1981), 1980US-0164074 (June 30, 1980)

http://westbrs:9000/bin/gate.exe?f=TOC&state=3pgbkl.3&ref=2&dbname=PGPB,USPT,US... 10/14/04

PATENT-FAMILY:

PUB-NO

PUB-DATE

LANGUAGE

PAGES

MAIN-IPC

US 4378347 A

March 29, 1983

005

INT-CL (IPC): A61K 35/55

ABSTRACTED-PUB-NO: US 4378347A

BASIC-ABSTRACT:

A dose of fibroblast growth factor (FGF) effective for treating the heart, suspended in a slow-release carrier for use in treatment of ischaemic heart disease is claimed. The carrier pref. comprises dextran beads. Admin. is pref. by direct injection into the heart, the dosage being 0.01-1000 mg per 100 g of heart. Intravenous, subcutaneous or oral admin. is also possible.

Admin. of the dose after myocardial infarction produces a sustained increase in blood flow in and around the damaged areas of myocardium and reduces the extent of damage.

Draw Desi	KWMC	Claims	0			eterence	Date	esification	0	Review	Front	Citation	Title	Full
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3. Document ID: US 4296100 A, GB 2090528 A, GB 2090528 B, JP 57500878 W, WO 8200098 A

L2: Entry 3 of 3

File: DWPI

Oct 20, 1981

DERWENT-ACC-NO: 1981-83435D

DERWENT-WEEK: 198145

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TITLE: Treating the heart with fibroblast growth factor - to alleviate conditions

caused by myocardial infarction(s)

INVENTOR: FRANCO, W P

PRIORITY-DATA: 1980US-0164074 (June 30, 1980), 1981US-0274722 (June 18, 1981)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
US 4296100 A	October 20, 1981		005	
GB 2090528 A	July 14, 1982		000	
GB 2090528 B	August 15, 1984		000	
JP 57500878 W	May 20, 1982		000	
WO 8200098 A	January 21, 1982	E	000	

INT-CL (IPC): A61K 9/00; A61K 35/55

ABSTRACTED-PUB-NO: GB 2090528B

BASIC-ABSTRACT:

Treatment of an area in the heart of a patient subjected to ischemic heart disease comprises admin. of fibroblast growth factor (FGF) to the heart.

After the treatment blood flow is increased for sustained periods after myocardial infarction. The treatment is useful after myocardial infarction (or when there is an indication of impeding myocardial infarction), when blood flow is increased in the treatment area and in surrounding areas. It is also useful with heart surgery procedures e.g. coronary by-pass operations, to reduce the quantity of myocardium

damage due to ischemic disease. FGF is a known mitogenic agent for a variety of mesodermal cells in vitro and it has been used to increase vascularisation in the cornea of laboratory animals. Dose is 10 micrograms-1g.100g heart by direct injection, esp. intravenously. The treatment is useful in man, cats, dogs, cows, etc.

ABSTRACTED-PUB-NO:

US 4296100A EQUIVALENT-ABSTRACTS:

A composition for use in the treatment of ischemic heart disease, comprising fibroblast growth factor (FGF) in association with a carrier material which ensures a slow release of the FGF therefrom, the carrier material being dextran or an albumin macro-aggregate.

Full	Title Citation	Front	Review	Classification	Date	Reference			Claims	KOME	Draw, Des
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	Terms						Documents				

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Search Results - Record(s) 1 through 4 of 4 returned.

☐ 1. Document ID: WO 2003033917 A1

Using default format because multiple data bases are involved.

L4: Entry 1 of 4

File: DWPI

Apr 24, 2003

DERWENT-ACC-NO: 2003-343371

DERWENT-WEEK: 200332

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: A double acting actuator for exerting tensile and thrust forces comprise a flexible annular cylinder with top and bottom end caps forming two chambers each with

an inlet for a pressurizing fluid

INVENTOR: FERRARESI, C; FRANCO, W; QUAGLIA, G

PRIORITY-DATA: 2001IT-T000984 (October 17, 2001)

PATENT-FAMILY:

PUB-NO

PUB-DATE

LANGUAGE

PAGES MAIN-IPC

WO 2003033917 A1

April 24, 2003

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017 F1

F15B015/10

INT-CL (IPC): $\underline{F15} \ \underline{B} \ \underline{15}/\underline{10}$

Full Title Citation Front Review Classification Date Reference Claims HWIC Draw. Desc

☐ 2. Document ID: IT 1292335 B

L4: Entry 2 of 4

File: DWPI

Jan 29, 1999

DERWENT-ACC-NO: 2001-467388

DERWENT-WEEK: 200151

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: Muscular actuator using fluid with straight fibers - NoAbstract

INVENTOR: FERRARESI, C; FRANCO, W; MANUELLO BERTETTO, A

PRIORITY-DATA: 1997IT-T000499 (June 9, 1997)

PATENT-FAMILY:

PUB-NO

PUB-DATE

LANGUAGE

PAGES

MAIN-IPC

IT 1292335 B

January 29, 1999

000

F15B000/00

INT-CL (IPC): $\underline{F15} \ \underline{B} \ \underline{0/00}$

☐ 3. Document ID: BE 1010130 A4

L4: Entry 3 of 4

File: DWPI

Jan 6, 1998

Jul 9, 1992

DERWENT-ACC-NO: 1998-077674

DERWENT-WEEK: 199808

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: Aeration of polluted earth for biological purification - using cutting and lifting rotors with efficient displacement for good aeration and subsequent

purification

INVENTOR: FRANCO, E; FRANCO, W

PRIORITY-DATA: 1996BE-0000265 (March 25, 1996)

PATENT-FAMILY:

PUB-NO

PUB-DATE

LANGUAGE

PAGES

MAIN-IPC

BE 1010130 A4

January 6, 1998

F

012

B09C000/00

INT-CL (IPC): $\underline{B09}$ \underline{C} $\underline{0/00}$; $\underline{C05}$ \underline{F} $\underline{0/00}$

ABSTRACTED-PUB-NO: BE 1010130A

BASIC-ABSTRACT:

A machine for aerating earth consists of a hollow structure, mounted on wheels, supporting adjacent and non-vertically mounted rotors. Also claimed is a method of treating earth using the machine described above, in particular to aerate for biological purification purposes, where the earth is treated in a long heap and redeposited behind the machine.

ADVANTAGE - The earth is displaced efficiently for good aeration and subsequent purification.

F	ull	Titl∈	Citation Front	Review	Classification	Date	Reference			Claims	KOMO	Draw, Des
***************************************	П	4	Document ID:	WO 9	211439 A1	I AI	9190804	A BE 100	04106 A3 F	EP 5619	36 A 1	

File: DWPI

DERWENT-ACC-NO: 1992-250145 DERWENT-WEEK: 199230

L4: Entry 4 of 4

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: Handling long elements such as pipes - involves upper frame carried by machine and connected to power take=off and intermediate frame which can be inclined to upper

frame

INVENTOR: FRANCO, E; FRANCO, W

PRIORITY-DATA: 1990BE-0001238 (December 20, 1990)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 WO 9211439 A1
 July 9, 1992
 E
 022
 E21B019/14

 AU 9190804 A
 July 22, 1992
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 E21B019/14

<u>BE 1004106 A3</u> September 22, 1992 F 016 E21B000/00 <u>EP 561936 A1</u> September 29, 1993 E 022 E21B019/14

INT-CL (IPC): E02F 3/96; E21B 19/14; E21B 19/15

ABSTRACTED-PUB-NO: WO 9211439A

BASIC-ABSTRACT:

The device for handling long elements has an upper frame (10) carried by a mobile machine (1) and an intermediate frame (20) is mounted on the upper frame. This intermediate frame can perform a movement of inclination with respect to the upper frame. In its turn the intermediate frame supports a lower frame (30) which can move in a longitudinal direction.

On the lower frame is a grip (40) capable of lifting the long elements and holding them. The power take off is a hydraulic power take-off.

USE/ADVANTAGE - Gives independent control to grip and hold long elements such as pipes.

Full Titl	e Citation f	Front Re	eview	Classificatio	n Dat	te Reference			Clai	ms l	KMC	Draw, Des
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Search Results - Record(s) 1 through 2 of 2 returned.

☐ 1. Document ID: US 20020058612 A1

Using default format because multiple data bases are involved.

L15: Entry 1 of 2

File: PGPB

May 16, 2002

PGPUB-DOCUMENT-NUMBER: 20020058612

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020058612 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor

and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: May 16, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

File: USPT

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/2; 424/43

	Full	Title	: Citation Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KAMC	Draw, Desi
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		2.	Document ID	: US 67	59386 B2							

US-PAT-NO: 6759386

L15: Entry 2 of 2

DOCUMENT-IDENTIFIER: US 6759386 B2

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor

and related proteins in the treatment of acute and chronic heart disease

DATE-ISSUED: July 6, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Jul 6, 2004

Franco; Wayne P.

Rocky Hill

CT

06067

US-CL-CURRENT: 514/2; 514/12, 514/14, 514/8, 530/300

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of <u>CPK-MB</u> levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart

http://westbrs:9000/bin/gate.exe?f=TOC&state=3pgbkl.16&ref=15&dbname=PGPB,USPT,U... 10/14/04

disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

24 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Ti	tle Citation Front Revie		Date Reference		Claims	KMC	Draw, Desi
Clear	Generate Collection	n Print	Fwd Refs	Bkwd Refs	Gener	ate O	ACS
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Hit List

Generate OACS Clear Generate Collection Print Fwd Refs **Bkwd Refs Search Results -** Record(s) 1 through 1 of 1 returned. ☐ 1. Document ID: US 6759386 B2 Using default format because multiple data bases are involved. L20: Entry 1 of 1 File: USPT Jul 6, 2004 US-PAT-NO: 6759386 DOCUMENT-IDENTIFIER: US 6759386 B2 TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease DATE-ISSUED: July 6, 2004 INVENTOR-INFORMATION: ZIP CODE COUNTRY CITY STATE NAME Franco; Wayne P. Rocky Hill CT06067 US-CL-CURRENT: 514/2; 514/12, 514/14, 514/8, 530/300 Full Title Citation Front Review Classification Date Reference Claims KMC Draw Des **Generate OACS** Clear Generate Collection Print Fwd Refs Bkwd Refs Terms Documents L19 AND CPK-MB Change Format Display Format: |-

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Generate OACS

Search Results - Record(s) 1 through 26 of 26 returned.

1. Document ID: US 20040185440 A9

Using default format because multiple data bases are involved.

L10: Entry 1 of 26

File: PGPB

Sep 23, 2004

PGPUB-DOCUMENT-NUMBER: 20040185440

PGPUB-FILING-TYPE: corrected

DOCUMENT-IDENTIFIER: US 20040185440 A9

TITLE: 125 human secreted proteins

PUBLICATION-DATE: September 23, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Feng, Ping	Gaithersburg	MD	US	
Ruben, Steven M.	Olney	MD	US	
Rosen, Craig A.	Laytonsville	MD	US	
Ebner, Reinhard	Gaithersburg	MD	US	
Olsen, Henrik S.	Gaithersburg	MD	US	
Ni, Jian	Rockville	MD .	US	
Wei, Ying-Fei	Berkeley	CA	US	
Soppet, Daniel R.	Centreville	VA	US	
Moore, Paul A.	Germantown	MD	US	
Kyaw, Hla	Frederick	MD	US	
LaFleur, David W.	Washington	DC	US	
Shi, Yanggu	Gaithersburg	MD	US	
Janat, Fouad	Westerly	RI	US	
Endress, Gregory A.	Potomac	MD	US	
Carter, Kenneth C.	North Potomac	MD	US	

US-CL-CURRENT: 435/6; 435/69.1, 514/2, 530/300, 536/23.1

Full	Citation	Front	Revieto	Classification	Cate	Reference	Sequences	Attachments	Claima	Emp).	[1]369
	Litation	Front	Memend	Classification	U3te	Reference	pedagunga	Augunmenti	Claime	- 111111	(113
ı		Citation	Citation Front	Citation Front Review	Citation Front Review Classification	Citation Front Review Classification Cate	Citation Front Review Classification Date Reference	Citation Front Review Classification Cate Reference Sequences	Citation Front Review Classification Date Reference Sequences Attachments	Citation Front Review Classification Date Reference Sequences Attachments Claims	Citation Front Review Classification Date Reference Sequences Attachments Claims 1990

2. Document 1D: US 2004009/401 A1

L10: Entry 2 of 26

File: PGPB

May 20, 2004

PGPUB-DOCUMENT-NUMBER: 20040097401

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040097401 A1

TITLE: Lysine in therapeutic angiogenesis, particularly in treating ischaemic conditions

PUBLICATION-DATE: May 20, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Datta, Debatosh

Kolkata

IN

US-CL-CURRENT: 514/2; 514/564, 514/565

ABSTRACT:

Present invention features methods for induction of angiogenesis by administration of lysine (l-&d-) or lysine oligomers (molecular weight approx between 500 and 2500), both homo and hetero-oligomers, consisting of either l-or d- or both enantiomers.

Induction of Angiogenesis by the methods of the invention can be use in therapeutic angiogenesis, in, for example, treatment of ischaemic conditions and syndromes, such as chronic wounds (e.g diabetic wounds and ulcers, bed sores and other pressure sores, burns of various degrees and extents etc.) as well as coronary and cerebral ischaemia and peripheral vascular ischaemic conditions. Induction of angiogenesis by the described methods also will be useful in inducing/enhancing radiosesitivity in some solid tumors.

l Tit	itl∈	Citation	Frent	Review	Classification	Date	Reference	Sequences	Attachments	Claims	1000	(rra
111	itle	Ultation	Front	Meinleim	Ulassmoation	Uate:	Meterence	pedagues.	Suscimence	Glaima	f. anti-	

3. Document ID: US 20040033971 A1

L10: Entry 3 of 26

File: PGPB

Feb 19, 2004

PGPUB-DOCUMENT-NUMBER: 20040033971

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040033971 A1

TITLE: Polypeptides and nucleic acids encoding same

PUBLICATION-DATE: February 19, 2004

NAME	CITY	STATE	COUNTRY	RULE-47
Gangolli, Esha A.	Madison	CT	US	
Patturajan, Meera	Branford	CT	US	
Vernet, Corine A.M.	Branford	CT	US	
Malyankar, Uriel M.	Branford	CT	US	
Kekuda, Ramesh	Norwalk	CT	US	
Stone, David J.	Guilford	CT	US	
Anderson, David	Branford	CT	US	

Nov 20, 2003

Shimkets, Richard A.	Guilford	CT	US
Burgess, Catherine E.	Wethersfield	CT	US
Zerhusen, Bryan D.	Branford	CT	US
Liu, Xiaohong	Branford	CT	US
Spytek, Kimberly A.	New Haven	CT	US
Casman, Stacie J.	North Haven	CT	US
Boldog, Ference L.	North Haven	CT	US
Smithson, Glennda	Guilford	CT	US
Li, Li	Branford	CT	US
Ji, Weizhen	Branford	CT	US
MacDougall, John R.	Hamden	CT	US

US-CL-CURRENT: 514/44; 435/320.1, 435/325, 435/6, 435/7.1, 514/2, 530/387.1, 536/23.1

ABSTRACT:

Disclosed herein are nucleic acid sequences that encode novel polypeptides. Also disclosed are polypeptides encoded by these nucleic acid sequences, and antibodies, which immunospecifically-bind to the polypeptide, as well as derivatives, variants, mutants, or fragments of the aforementioned polypeptide, polynucleotide, or antibody. The invention farther discloses therapeutic, diagnostic and research methods for diagnosis, treatment, and prevention of disorders involving any one of these novel human nucleic acids and proteins.

Full	Titl∈	Citation	Front	Review	Classification	Late	Reference	Sequences	###achments	Claims	15000	Draw Do
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File: PGPB

PGPUB-DOCUMENT-NUMBER: 20030215840

PGPUB-FILING-TYPE: new

L10: Entry 4 of 26

DOCUMENT-IDENTIFIER: US 20030215840 A1

TITLE: Methods and compositions for treating cardiovascular disease using 1682, 6169, 6193, 7771, 14395, 29002, 33216, 43726, 69292, 26156, 32427, 2402, 7747, 1720, 9151, 60491, 1371, 7077, 33207, 1419, 18036, 16105, 38650, 14245, 58848, 1870, 25856, 32394, 3484, 345, 9252, 9135, 10532, 18610, 8165, 2448, 2445, 64624, 84237, 8912, 2868, 283, 2554, 9464, 17799, 26686, 43848, 32135, 12208, 2914, 51130, 19489, 21833, 2917, 59590, 15992, 2094, 2252, 3474, 9792, 15400, 1452 or 6585 molecules

PUBLICATION-DATE: November 20, 2003

NAME	CITY	STATE	COUNTRY	RULE-47
Logan, Thomas J.	Springfield	PA	US	
Chun, Miyoung	Belmont	MA	US	
Galvin, Katherine M.	Jamaica Plain	MA	US	
Healy, Aileen	Medford	MA	US	

Acton, Susan L.	Lexington	MA	US
Donoghue, Mary A.	West Roxbury	MA	US
Stagliano, Nancy	North Reading	MA	US
Perodin, Jacqueline	Arlington	MA	US
Rodrigue-Way, Amelie	Malden	MA	US

US-CL-CURRENT: 435/6; 424/146.1, 435/7.2, 514/1, 514/2, 514/44

ABSTRACT:

The present invention relates to methods for the diagnosis and treatment of cardiovascular disease, including, but not limited to, atherosclerosis, reperfusion injury, hypertension, restenosis, arterial inflammation, heart failure, thrombosis and endothelial cell disorders. Specifically, the present invention identifies the differential expression of 1682, 6169, 6193, 7771, 14395, 29002, 33216, 43726, 69292, 21656, 32427, 2402, 7747, 1720, 9151, 60491, 1371, 7077, 33207, 1419, 18036, 16105, 38650, 14245, 58848, 1870, 25856, 32394, 3484, 345, 9252, 9135, 10532, 18610, 8165, 2448, 2445, 64624, 84237, 8912, 2868, 283, 2554, 9464, 17799, 26686, 43848, 32135, 12208, 2914, 51130, 19489, 21833, 2917, 59590, 15992, 2094, 2252, 3474, 9792, 15400, 1452 and 6585 genes in cardiovascular disease states, relative to their expression in normal, or non-cardiovascular disease states, and/or in response to manipulations relevant to cardiovascular disease. The present invention describes methods for the diagnostic evaluation and prognosis of various cardiovascular diseases, and for the identification of subjects exhibiting a predisposition to such conditions. The invention also provides methods for identifying a compound capable of modulating cardiovascular disease. The present invention also provides methods for the identification and therapeutic use of compounds as treatments of cardiovascular disease.

Full Title Citation Front Review Classification	Date Refere	nce Sequences	Attachments	Claims	Philips	िएअल्स् (ल
☐ 5. Document ID: US 20030215452	2 A1					
L10: Entry 5 of 26	1 7	PGPB		Nov	20	2002

PGPUB-DOCUMENT-NUMBER: 20030215452

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030215452 A1

TITLE: Methods and compositions for treating hematological disorders using 131, 148, 199, 12303, 13906, 15513, 17822, 302, 5677, 194, 14393, 28059, 7366, 12212, 1981, 261, 12416, 270, 1410, 137, 1871, 13051, 1847, 1849, 15402, 340, 10217, 837, 1761, 8990 or 13249 molecules

PUBLICATION-DATE: November 20, 2003

NAME	CITY	STATE	COUNTRY	RULE-47
Carroll, Joseph M.	Cambridge	MA	US	
Healy, Aileen	Medford	MA	US	
Weich, Nadine S.	Brookline	MA	US	
Kelly, Louise M.	Brookline	MA	US	

US-CL-CURRENT: 424/146.1; 435/6, 435/7.2, 514/1, 514/2, 514/44

ABSTRACT:

The present invention relates to methods for the diagnosis and treatment of hematological disorders. Specifically, the present invention identifies the differential expression of 131, 148, 199, 12303, 13906, 15513, 17822, 302, 5677, 194, 14393, 28059, 7366, 12212, 1981, 261, 12416, 270, 1410, 137, 1871, 13051, 1847, 1849, 15402, 340, 10217, 837, 1761, 8990 and 13249 genes in tissues relating to hematological disorders sensation, relative to their expression in normal, or non-hematological disorders disease states, and/or in response to manipulations relevant to hematological disorders. The present invention describes methods for the diagnostic evaluation and prognosis of various hematological disorders, and for the identification of subjects exhibiting a predisposition to such conditions. The invention also provides methods for identifying a compound capable of modulating hematological disorders. The present invention also provides methods for the identification and therapeutic use of compounds as treatments of hematological disorders.

Full	Title	Citation	Feent	Review	Classitication	Crate	Flataran	sa Sequenc	es Attachments	Claims	Finde	Станц Со
	6.]	Docume	nt ID:	US 200	3021147	2 A1						
L10:	Entry	/ 6 of	26				File:	PGPB		Nov	13,	2003

PGPUB-DOCUMENT-NUMBER: 20030211472

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030211472 A1

TITLE: 125 human secreted proteins

PUBLICATION-DATE: November 13, 2003

THE DISTORCE THE CHARLES TO THE				
NAME	CITY	STATE	COUNTRY	RULE-47
Feng, Ping	Gaithersburg	MD	US	
Ruben, Steven M.	Olney	MD	US	
Rosen, Craig A.	Laytonsville	MD	US	
Ebner, Reinhard	Gaithersburg	MD	US	
Olsen, Henrik S.	Gaithersburg	MD	US	
Ni, Jian	Rockville	MD	US	
Wei, Ying-Fei	Berkeley	CA	US	
Soppet, Daniel R.	Centreville	VA	US	
Moore, Paul A.	Germantown	MD	US	
Kyaw, Hla	Frederick	MD	US	
LaFleur, David W.	Washington	DC	US	
Shi, Yanggu	Gaithersburg	MD	US	
Janat, Fouad	Westerly	RI	US	
Endress, Gregory A.	Potomac	MD	US	
Carter, Kenneth C.	North Potomac	MD	US	

US-CL-CURRENT: 435/6; 435/69.1, 514/2, 530/300, 536/23.1

ABSTRACT:

The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins.

Full	Title Citation	Front	Flevieu	Classification	Date	Reference	Sequences	attachments	Claims	15640	Erass D
			X 7 0 0 0	020100425						***************************************	
	7. Docume	ent ID:	US 20	030199425	AI						
L10:	Entry 7 of	26				File: P	GPB		Oct	23,	2003

PGPUB-DOCUMENT-NUMBER: 20030199425

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030199425 A1

TITLE: Compositions and methods for treatment of hyperplasia

PUBLICATION-DATE: October 23, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47
Desai, Neil P. Los Angeles CA US
Soon-Shiong, Patrick Los Angeles CA US

US-CL-CURRENT: 514/2; 424/45, 514/291, 514/365, 514/449

ABSTRACT:

In accordance with the present invention, there are provided methods for treating hyperplasia in a subject in need thereof. In another aspect of the invention, there are provided methods for reducing neointimal hyperplasia associated with vascular interventional procedures. Formulations contemplated for use herein comprise proteins and at least one pharmaceutically active agent.

Full	Titl∈	Citation	Front	Review	Classification	Date	Reference	Зедиелюев	Attachmenta	Claimz	[huli]	Draws D
	8.	Docume	nt ID:	US 20	030154504	A1						

PGPUB-DOCUMENT-NUMBER: 20030154504

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030154504 A1

TITLE: Methods and compositions for modulating carbohydrate metabolism

PUBLICATION-DATE: August 14, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Farese, Robert V. JR. San Francisco CA US Chen, Hubert C. San Francisco CA US

US-CL-CURRENT: 800/18; 514/2, 514/3

ABSTRACT:

Methods and compositions for modulating carbohydrate metabolism in a host are provided. In the subject methods, diacylglycerol acyltransferase (DGAT) activity (specifically DGAT1 activity) is modulated, e.g., reduced or enhanced, to achieve a desired insulin and/or leptin sensitivity, thereby modulating carbohydrate metabolism, e.g., increasing or decreasing blood glucose levels, glucose uptake into cells and assimilation into glycogen. Also provided are pharmaceutical compositions for practicing the subject methods. The subject methods and compositions find use in a variety of applications, including the treatment of hosts suffering conditions associated with abnormal carbohydrate metabolism, such as obesity or diabetes.

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Full T	Title	Citation	Frent	Register	Classification	trate	NeTerebor	Dediciplines	Sugnification	LIBINI	E. 3855.6575	F. Leton.

9. Document ID: US 20030152574 A1

L10: Entry 9 of 26 File: PGPB Aug 14, 2003

PGPUB-DOCUMENT-NUMBER: 20030152574

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030152574 A1

TITLE: Methods and compositions to treat cardiovascular disease using 1419, 58765 and 2210

PUBLICATION-DATE: August 14, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Logan, Thomas Joseph Springfield PA US Chun, Miyoung Belmont MA US

US-CL-CURRENT: 424/146.1; 435/7.2, 514/1, 514/2, 514/44

ABSTRACT:

The present invention relates to methods for the diagnosis and treatment of cardiovascular disease, including, but not limited to, atherosclerosis, reperfusion injury, hypertension, restenosis, arterial inflammation, thrombosis and endothelial cell disorders. Specifically, the present invention identifies the differential

expression of 1419, 58765 or 2210 genes in cardiovascular disease states, relative to their expression in normal, or non-cardiovascular disease states, and/or in response to manipulations relevant to cardiovascular disease. The present invention describes methods for the diagnostic evaluation and prognosis of various cardiovascular diseases, and for the identification of subjects exhibiting a predisposition to such conditions. The invention also provides methods for identifying a compound capable of modulating cardiovascular disease. The present invention also provides methods for the identification and therapeutic use of compounds as treatments of cardiovascular disease.

Full	Title	Citation	Front	Fleview	Classification	Date	Reference	. Sequences	:#ttaclments	Claims	15000	[it3bit [i-
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	10.	Docum	ent ID): US 2	003009265	8 A1						
L10:	Entr	y 10 of	26				File:	PGPB		May	15,	2003

PGPUB-DOCUMENT-NUMBER: 20030092658

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030092658 A1

TITLE: Novel human enzyme family members and uses thereof

PUBLICATION-DATE: May 15, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Meyers, Rachel E.	Newton	MA	US	
Glucksmann, Maria Alexandra	Lexington	MA	US	
Rudolph-Owen, Laura A.	Jamaica Plain	MA	US	

US-CL-CURRENT: 514/44; 424/130.1, 435/6, 514/2

ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 33312, 33303, 32579, 21509, 33770, 46638, and 50090 nucleic acid molecules, which encode novel G protein-coupled receptor family members, human thioredoxin family members, human leucine-rich repeat family members, and human ringfinger family member. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 33312, 33303, 32579, 21509, 33770, 46638, or 50090 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 33312, 33303, 32579, 21509, 33770, 46638, or 50090 gene has been introduced or disrupted. The invention still further provides isolated 33312, 33303, 32579, 21509, 33770, 46638, or 50090 proteins, fusion proteins, antigenic peptides and anti-33312, 33303, 32579, 21509, 33770, 46638, or 50090 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

11. Document ID: US 20030083231 A1

L10: Entry 11 of 26

File: PGPB

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030083231

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030083231 A1

TITLE: Blood cell deficiency treatment method

PUBLICATION-DATE: May 1, 2003

INVENTOR - INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Ahlem, Clarence N.	San Diego	CA	US	
Reading, Christopher	San Diego	CA	US	
Frincke, James	San Diego	CA	ບຣ	
Stickney, Dwight	Granite Bay	CA	us	
Lardy, Henry A.	Madison	WI	US	
Marwah, Padma	Middleton	WI	us	
Marwah, Ashok	Middleton	WI	US	
Prendergast, Patrick T.	Straffan		IE	

US-CL-CURRENT: 514/2; 514/169, 514/173, 514/26, 514/44, 514/63

ABSTRACT:

The invention relates to the use of compounds to treat a number of conditions, such as thrombocytopenia, neutropenia or the delayed effects of radiation therapy. Compounds that can be used in the invention include methyl-2,3,4-trihydroxy-1-O-(7,17-dioxoandrost-5-ene-3.beta.-yl)-.beta.-D--glucopyranosiduronate, 16.alpha.,3.alpha.-dihydroxy-5.alpha.-androstan-17--one or 3,7,16,17-tetrahydroxyandrost-5-ene, 3,7,16,17-tetrahydroxyandrost--4-ene,3,7,16,17-tetrahydroxyandrost--4-ene,3,7,16,17-tetrahydroxyandrost- tane that can be used in the treatment method.

tie (Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	10000	[гане, [
į l		Citation	Citation Front	Citation Front Review	Citation Front Review Classification	Citation Front Review Classification Date	Citation Front Review Classification Date Reference	Citation Front Review Classification Date Reference Sequences	Citation Front Review Classification Date Reference Sequences Attachments	Citation Front Review Classification Date Reference Sequences Attachments Claims	Citation Front Review Classification Date Reference Sequences Attachments Claims 1990

12. Document ID: US 20030073118 A1

L10: Entry 12 of 26

File: PGPB

Apr 17, 2003

PGPUB-DOCUMENT-NUMBER: 20030073118

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030073118 A1

TITLE: MID 9002, a human sulfatase family member and uses therefor

PUBLICATION-DATE: April 17, 2003

NAME

CITY

STATE COUNTRY

RULE-47

Williamson, Mark W.

Saugus MA

US

US-CL-CURRENT: 435/6; 424/130.1, 514/1, 514/2, 514/44

ABSTRACT:

The invention provides isolated nucleic acids molecules, designated MID 9002 nucleic acid molecules, which encode novel sulfatase family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing MID 9002 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a MID 9002 gene has been introduced or disrupted. The invention still further provides isolated MID 9002 proteins, fusion proteins, antigenic peptides and anti-MID 9002 antibodies. Diagnostic and therapeutic methods utilizing compositions of the invention are also provided.

Full	Titl∈	Citation	Front	Flevien	Classification	Date	Reference	. Sequences	Attachments	Claims	E)0040	frant fe
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	13.	Docum	ent ID	: US 2	002015104	6 A1						
L10:	Entr	y 13 of	26				File:	PGPB		Oct	17,	2002

PGPUB-DOCUMENT-NUMBER: 20020151046

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020151046 A1

TITLE: 52871, a novel human G protein coupled receptor and uses thereof

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Glucksmann, Maria Alexandra Lexington MA US

Glucksmann, Maria Alexandra Lexington MA US Silos-Santiago, Inmaculada Cambridge MA US

US-CL-CURRENT: 435/320.1; 435/325, 435/6, 435/69.1, 435/7.1, 514/2, 530/324, 530/387.7, 536/23.5

ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 52871 nucleic acid molecules, which encode novel G-Protein Coupled Receptor molecules. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 52871 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 52871 gene has been introduced or disrupted. The invention still further provides isolated 52871 proteins, fusion proteins, antigenic peptides and anti-52871 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.



14. Document ID: US 20020119913 A1

L10: Entry 14 of 26

File: PGPB

Aug 29, 2002

May 23, 2002

PGPUB-DOCUMENT-NUMBER: 20020119913

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020119913 A1

TITLE: 61833, a novel human pyridoxyl-dependent decarboxylase family member and

uses thereof

PUBLICATION-DATE: August 29, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Glucksmann, Maria Alexandra Lexington MA US

US-CL-CURRENT: 514/2; 435/320.1, 435/325, 435/6, 435/69.1, 435/7.2, 530/324,

530/387.9, 536/23.5

ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 61833 nucleic acid molecules, which encode novel pyridoxyl-dependent decarboxylase members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 61833 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 61833 gene has been introduced or disrupted. The invention still further provides isolated 61833 proteins, fusion proteins, antigenic peptides and anti-61833 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

Full	Title	Citation From	t Review	Classification	Cate	Reference	Sequences	Attachments	Claime	15040	Dram D
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File: PGPB

PGPUB-DOCUMENT-NUMBER: 20020061521

PGPUB-FILING-TYPE: new

L10: Entry 15 of 26

DOCUMENT-IDENTIFIER: US 20020061521 A1

TITLE: Nucleic acids, proteins, and antibodies

PUBLICATION-DATE: May 23, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Rosen, Craig A. Laytonsville MD US

Ruben, Steven M.

Olney

MD

US

Barash, Steven C.

Rockville

MD

US

US-CL-CURRENT: 435/6; 435/69.1, 514/2, 530/300, 536/23.1

ABSTRACT:

The present invention relates to novel cardiovascular system related polynucleotides and the polypeptides encoded by these polynucleotides herein collectively known as "cardiovascular system antigens," and the use of such cardiovascular system antigens for detecting disorders of the cardiovascular system, particularly the presence of cancer of cardiovascular system tissues and cancer metastases. More specifically, isolated cardiovascular system associated nucleic acid molecules are provided encoding novel cardiovascular system associated polypeptides. Novel cardiovascular system polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human cardiovascular system associated polynucleotides and/or polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to the cardiovascular system, including cancer of cardiovascular system tissues, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting the production and function of the polypeptides of the present invention.

Full	Titl≥	Citation F	iont	Review	Classification	Date	Reference	Sequences	Attachments	Claims	1,30000	francti-
	16	Docume	nt ID:	US 2	002005861	2 A1						
		y 16 of					File:	PGPB		May	16,	2002

PGPUB-DOCUMENT-NUMBER: 20020058612

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020058612 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: May 16, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Franco, Wayne P. Rocky Hill CT US

US-CL-CURRENT: 514/2; 424/43

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment

with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

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17. Document ID: US 20020037832 A1

L10: Entry 17 of 26

File: PGPB

Mar 28, 2002

PGPUB-DOCUMENT-NUMBER: 20020037832

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020037832 A1

TITLE: Use of alpha-MSH and EPO for preventing or treating ischemic conditions

PUBLICATION-DATE: March 28, 2002

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47
Nielsen, Soren Abyhoj DK
Frokiaer, Jorgen Abyhoj DK
Jonassen, Thomas Engelbrecht Norkild Frederiksberg DK
Bjerke, Thorbjorn Fredensborg DK

US-CL-CURRENT: 514/2; 514/169

ABSTRACT:

Alpha--melanocyte stimulating hormone (.alpha.-MSH) or an equivalent is used, in conjunction with erythropoietin (EPO) or equivalent, to prevent or treat ischemic conditions.

Full	Titl∈	Citation	Front	Review	Classification	Cate	Reference	Sequences	Attachments	Claims	15640	Grand C
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	10	Docum	ent ID	· IIS 6	787519 B2							
	18.	Docum		. 000	1101212							

US-PAT-NO: 6787519

DOCUMENT-IDENTIFIER: US 6787519 B2

TITLE: Methods of treating disorders related to apoE

DATE-ISSUED: September 7, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Huang; Yadong San Francisco CA

Record List Display

Mahley; Robert W.

San Francisco

ÇA

US-CL-CURRENT: 514/2; 514/17, 514/18, 530/300, 530/329

ABSTRACT:

The present invention provides methods inhibiting formation of neurofibrillary tangles; and methods for treating disorders relating to apolipoprotein E (apoE) in a subject. The methods generally involve reducing the level of a carboxyl-terminal truncated form of apoE in a neuronal cell of a subject. The invention further provides isolated cells comprising a nucleic acid molecule encoding a carboxylterminal truncated form of apoE; and methods of screening compounds using the cells. The invention further provides compounds that inhibit an apoE cleavage enzyme, and that reduce the formation of neurofibrillary tangles in a neuronal cell. The invention further provides transgenic non-human animals that include as a transgene a nucleic acid that encodes a carboxyl-terminal truncated form of apoE; as well as methods of screening compounds using transgenic animals.

16 Claims, 15 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 8

Full	Titl∈	Citation	Foont	Review	Classinication	Crate	Reference	Was Walle	Claims	Polic	Cram Cr
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	19.	Docum	ent ID): US 6	759386 B2						
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US-PAT-NO: 6759386

L10: Entry 19 of 26

DOCUMENT-IDENTIFIER: US 6759386 B2

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

DATE-ISSUED: July 6, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE.

ZIP CODE

COUNTRY

Franco; Wayne P.

Rocky Hill

CT

06067

US-CL-CURRENT: 514/2; 514/12, 514/14, 514/8, 530/300

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

24 Claims, 4 Drawing figures

Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Claims 1960 Prace De Full Title Citation Front Review Classification Date Reference 20. Document ID: US 6737404 B2 May 18, 2004 L10: Entry 20 of 26 File: USPT

US-PAT-NO: 6737404

DOCUMENT-IDENTIFIER: US 6737404 B2

TITLE: Methods of using analogs of human basic fibroblast growth factor mutated at

one or more of the positions glutamate 89, aspartate 101 or leucine 137

DATE-ISSUED: May 18, 2004

INVENTOR-INFORMATION:

ZIP CODE COUNTRY STATE CITY NAME Wilmington DE Springer; Barry A.

Boxford PA Pantoliano; Michael W. Sharp; Celia M. Doylestown PA

US-CL-CURRENT: 514/12; 514/2, 530/399

ABSTRACT:

The present invention relates to novel muteins of human basic fibroblast growth factor with superagonist properties. Both protein and the respective encoding nucleic acid species are disclosed. The invention also embodies vectors and host cells for the propagation of said nucleic acid sequences and the production of said muteins. Also disclosed are methods for stimulating cell division, treating a wound, treating ischemia, treating heart disease, treating neural injury, treating peripheral vascular disease, treating a gastric ulcer and treating a duodenal ulcer.

30 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full	Titl∈	Citation	Front	Review	Classification	Crate	Reference		. Committee of the comm	Claims	1000	[rram [re
	21.	Docume	nt ID	: US 6	605592 B2							
L10:	Entr	y 21 of	26				File:	USPT		Aug	12,	2003

US-PAT-NO: 6605592

DOCUMENT-IDENTIFIER: US 6605592 B2

TITLE: Protein HOFNF53

DATE-ISSUED: August 12, 2003

INVENTOR-INFORMATION:					
NAME	CITY	STATE	ZIP	CODE	COUNTRY
Ni; Jian	Germantown	MD			
Baker; Kevin P.	Darnestown	MD			
Birse; Charles E.	North Potomac	MD			
Ebner; Reinhard	Gaithersburg	MD			
Fiscella; Michele	Bethesda	MD			
Komatsoulis; George A.	Silver Spring	MD			
LaFleur; David W.	Washington	DC			
Moore; Paul A.	Germantown	MD			
Olsen; Henrik S.	Gaithersburg	MD			
Rosen; Craig A.	Laytonsville	MD			
Ruben; Steven M.	Olney	MD			
Soppet; Daniel R.	Centreville	VA			
Young; Paul E.	Gaithersburg	MD			
Wei; Ping	Brookeville	MD			
Florence; Kimberly A.	Rockville	MD			

US-CL-CURRENT: 514/2; 435/252.3, 435/254.11, 435/320.1, 435/325, 435/471, 435/69.1, 435/71.1, 435/71.2, 514/12, 514/8, 530/350

ABSTRACT:

The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. In particular, the present application relates to a novel human protein, Protein HOFNF53. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

19 Claims, 22 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 22

Full	Title	Citation	Front	Review	Classification	Late	Reference		PT. FLET	Claima	Finds)	Fram. D
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	22.	Docum	ent ID): US 6	541224 B2							
L10:	Entry	/ 22 of	26				File:	USPT		Apr	1,	2003

US-PAT-NO: 6541224

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DOCUMENT-IDENTIFIER: US 6541224 B2

** See image for Certificate of Correction **

TITLE: Tumor necrosis factor delta polypeptides

DATE-ISSUED: April 1, 2003

INVENTOR-INFORMATION:

COUNTRY ZIP CODE STATE CITY

Berkeley CA Yu; Guo-Liang Germantown MD

Ni; Jian MD Rockville Gentz; Reiner L. CA

Carlsbad Dillon; Patrick J.

US-CL-CURRENT: 435/69.5; 435/69.1, 435/69.7, 435/7.71, 435/70.1, 514/12, 514/2, 530/350, 530/351

ABSTRACT:

The invention relates to human TNF delta and TNF epsilon polypeptides, polynucleotides encoding the polypeptides, methods for producing the polypeptides, in particular by expressing the polynucleotides, and agonists and antagonists of the polypeptides. The invention further relates to methods for utilizing such polynucleotides, polypeptides, agonists and antagonists for applications, which relate, in part, to research, diagnostic and clinical arts.

50 Claims, 7 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 11

Full	Titl∈	Citation	Front	Review	Classification	Cate	Reference			Claims	15640	Errand
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23. Document ID: US 6521211 B1

L10: Entry 23 of 26

File: USPT

Feb 18, 2003

US-PAT-NO: 6521211

DOCUMENT-IDENTIFIER: US 6521211 B1

TITLE: Methods of imaging and treatment with targeted compositions

DATE-ISSUED: February 18, 2003

INVENTOR-INFORMATION:

COUNTRY ZIP CODE STATE CITY NAME

A 7. Tucson Unger; Evan C. Tucson AZWu; Yunqiu

US-CL-CURRENT: 424/9.52; 424/450, 424/9.5, 424/9.51, 514/18, 514/2, 600/431, 600/437

ABSTRACT:

Novel ultrasound methods comprising administering to a patient a targeted vesicle composition which comprises vesicles comprising a lipid, protein or polymer, encapsulating a gas, in combination with a targeting ligand, and scanning the patient using ultrasound. The scanning may comprise exposing the patient to a first type of ultrasound energy and then interrogating the patient using a second type of ultrasound energy. The targeting ligand preferably targets tissues, cells or

receptors, including myocardial cells, endothelial cells, epithelial cells, tumor cells and the glycoprotein GPIIbIIIa receptor. The methods may be used to detect a thrombus, enhancement of an old or echogenic thrombus, low concentrations of vesicles and vesicles targeted to tissues, cells or receptors.

58 Claims, 17 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 12

Full Title Citation Front Review Clarafication Crate Reference Claims 1000 Prass D 24. Document ID: US 6475796 B1 Nov 5, 2002 L10: Entry 24 of 26 File: USPT

US-PAT-NO: 6475796

DOCUMENT-IDENTIFIER: US 6475796 B1

TITLE: Vascular endothelial growth factor variants

DATE-ISSUED: November 5, 2002

INVENTOR-INFORMATION:

ZIP CODE COUNTRY CITY STATE NAME

Los Altos Pollitt; N. Stephen San Jose CA Abraham; Judith A.

US-CL-CURRENT: 435/455; 424/198.1, 514/2, 530/350

ABSTRACT:

The invention is directed to a method of enhancing the biological activity of vascular endothelial growth factors (VEGF). The invention further concerns certain VEGF variants having enhanced biological activity, methods and means for preparing these variants, and pharmaceutical compositions comprising them. In a further aspect, the invention concerns methods of treatment using, and articles of manufacture containing such VEGF variants.

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17 Claims, 17 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 17

Full	Titl≟	Citation	Front	Review	Classification	Cate	Reference			Claims	150000	[11300
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US-PAT-NO: 6407135

L10: Entry 25 of 26

DOCUMENT-IDENTIFIER: US 6407135 B1

Jun 11, 2002

TITLE: Conjugates of dithiocarbamates with pharmacologically active agents and uses therefor

DATE-ISSUED: June 18, 2002

INVENTOR - INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Lai; Ching-San Encinitas CA Wang; Tingmin San Marcos CA

US-CL-CURRENT: 514/423; 514/2, 514/514, 530/402, 548/565, 548/573

ABSTRACT:

In accordance with the present invention, there are provided conjugates of nitric oxide scavengers (e.g., dithiocarbamates, or "DC") and pharmacologically active agents (e.g., NSAIDs). Invention conjugates provide a new class of pharmacologically active agents (e.g., anti-inflammatory agents) which cause a much lower incidence of side-effects due to the protective effects imparted by modifying the pharmacologically active agents as described herein. In addition, invention conjugates are more effective than unmodified pharmacologically active agents because cells and tissues contacted by the pharmacologically active agent(s) are protected from the potentially damaging effects of nitric oxide overproduction induced thereby as a result of the co-production of nitric oxide scavenger (e.g., dithiocarbamate), in addition to free pharmacologically active agent, when invention conjugate is cleaved.

21 Claims, 5 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 5

Full	Titl∈	Citation Fron	it Review	Classification	Late	Reference	**************************************	Claime	15000	[stand [
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File: USPT

US-PAT-NO: 6403552

L10: Entry 26 of 26

DOCUMENT-IDENTIFIER: US 6403552 B1

TITLE: Ob receptor and methods for the diagnosis and treatment of body weight disorders

DATE-ISSUED: June 11, 2002

INVENTOR-INFORMATION:

COUNTRY STATE ZIP CODE NAME CITY Watertown MA Tartaglia; Louis A. MA Weston Tepper; Robert I. Culpepper; Janice A. Brookline MΑ Holbrook MA White; David W.

US-CL-CURRENT: 514/2; 424/143.1, 435/69.7, 536/23.4

ABSTRACT:

The present invention relates to the discovery, identification and characterization of nucleotides that encode Ob receptor (ObR), a receptor protein that participates in mammalian body weight regulation. The invention encompasses obR nucleotides, host cell expression systems, ObR proteins, fusion proteins, polypeptides and peptides, antibodies to the receptor, transgenic animals that express an obR transgene, or recombinant knock-out animals that do not express the ObR, antagonists and agonists of the receptor, and other compounds that modulate obR gene expression or ObR activity that can be used for diagnosis, drug screening, clinical trial monitoring, and/or the treatment of body weight disorders, including but not limited to obesity, cachexia and anorexia.

41 Claims, 40 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 34

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1. Document ID: US 20040167070 A1

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L20: Entry 1 of 8

File: PGPB

Aug 26, 2004

PGPUB-DOCUMENT-NUMBER: 20040167070

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040167070 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: August 26, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

Full Title Citation Front Review Classificat	ion Date Reference	Sequences	#ttachments	Claims	150000	Citabe Ci-
☐ 2. Document ID: US 200401163	349 A1					
L20: Entry 2 of 8	File: PG	PB		Jun	17,	2004

PGPUB-DOCUMENT-NUMBER: 20040116349

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040116349 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: June 17, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full Title Citation Front Review Classification Date Reference Sequence: Attachments Claims Foot Grand December 2. Attachments Claims Foot Grand December 2.

PGPUB-DOCUMENT-NUMBER: 20040023863

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040023863 A1

TITLE: Methods of use growth factors for treating heart disease

PUBLICATION-DATE: February 5, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Franco, Wayne P.

Rocky Hill

CT

US

US-CL-CURRENT: 514/12

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral <u>inhalation therapy</u>.

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4. Document ID: US 20030036773 A1

L20: Entry 4 of 8

File: PGPB

Feb 20, 2003

PGPUB-DOCUMENT-NUMBER: 20030036773

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030036773 A1

TITLE: Systems and methods for treatment of coronary artery disease

PUBLICATION-DATE: February 20, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Whitehurst, Todd K.	Frazier Park	CA	US	
McGivern, James P.	Stevenson Ranch	CA	US	
McClure, Kelly H.	Simi Valley	CA	US	
Stultz, Mark R.	Maple Grove	MN	US	

US-CL-CURRENT: 607/3; 607/120

ABSTRACT:

Methods and systems for treatment of coronary artery disease (CAD) include implantation of the discharge portion(s) of a catheter and, optionally, electrodes on a lead, near the tissue(s) to be stimulated. Stimulation pulses, i.e., drug infusion pulses and optional electrical pulses, are supplied by a stimulator implanted remotely, and through the catheter or lead, which is tunneled subcutaneously between the stimulator and stimulation site. Stimulation sites include the coronary arteries, the aorta, the left ventricle, the left atrium, and/or the pulmonary veins, among other locations. Disclosed treatments include drugs used for acute treatment of CAD, for chronic treatment of CAD, to promote angiogenesis, and/or as AGE Crosslink Breakers, among other drugs. For instance, the systems and methods reduce or eliminate the incidence of CAD and related morbidities, improve symptoms resulting from CAD, and/or improve cardiac blood flow, cardiac function, and patient quality of life.

Full	Title	Citation	Frent	Review	Classification	Date	Referenc	e Sequencesi	.4ttachments	Claima	LindC	Frank Fr
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	5.]	Documer	nt ID:	US 20	020103454	A1						
L20:	Entry	y 5 of	8				File:	PGPB		Aug	1,	2002

PGPUB-DOCUMENT-NUMBER: 20020103454

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020103454 A1

TITLE: External addition of pulses to fluid channels of body to release or suppress endothelial mediators and to determine effectiveness of such intervention

PUBLICATION-DATE: August 1, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47
Sackner, Marvin A. Miami Beach FL US
Inman, D. Michael Miami FL US

US-CL-CURRENT: 604/19

ABSTRACT:

Methods of medical treatment and diagnosis using mediators released by endothelial cells stimulated by external addition of pulses to the circulation are disclosed. The external pulses produce circumferential shear stress in body fluid channels

that subsequently stimulates the endothelial cells to produce mediators that become available for therapeutic and diagnostic purposes. The preferred means of adding external pulses is the mechanical inducement of periodic acceleration of the body or parts of the body by a reciprocating motion platform.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims Doub Irraw De Grant De Grant

PGPUB-DOCUMENT-NUMBER: 20020058612

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020058612 A1

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

PUBLICATION-DATE: May 16, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Franco, Wayne P. Rocky Hill CT US

US-CL-CURRENT: 514/2; 424/43

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

Full	Title Citation Front	Review Classification	Date	Reference	Sequences	Lattachments	Claims	15040	[-[360 [6
П	7. Document ID:	US 6759386 B2							
T.20 ·	Entry 7 of 8			File: U	SPT		Jul	6,	2004

US-PAT-NO: 6759386

DOCUMENT-IDENTIFIER: US 6759386 B2

TITLE: Methods of use of fibroblast growth factor, vascular endothelial growth factor and related proteins in the treatment of acute and chronic heart disease

DATE-ISSUED: July 6, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Franco; Wayne P.

Rocky Hill

CT

06067

US-CL-CURRENT: 514/2; 514/12, 514/14, 514/8, 530/300

ABSTRACT:

Disclosed herein is a rational, multi-tier approach to the administration of growth factor proteins in the treatment of heart disease. Also disclosed is a method to evaluate the effectiveness of the administration of growth factor proteins comprising the clinical assay of CPK-MB levels in a patient undergoing treatment with growth factor proteins. In addition, there is disclosed a method for treatment of heart disease comprising administration of a therapeutically effective amount of a growth factor protein by oral inhalation therapy.

24 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4



☐ 8. Document ID: US 6673908 B1

L20: Entry 8 of 8

File: USPT

Jan 6, 2004

US-PAT-NO: 6673908

DOCUMENT-IDENTIFIER: US 6673908 B1

TITLE: Tumor necrosis factor receptor 2

DATE-ISSUED: January 6, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Stanton, Jr.; Vincent P.

Belmont

MΑ

US-CL-CURRENT: 536/22.1; 435/6, 435/91.1, 435/91.2, 536/23.1, 536/24.3, 536/24.31, 536/24.33

ABSTRACT:

The present disclosure describes the use of genetic variance information for genes involved in inflammatory or immunologic disease, disorder, or dysfunction. The variance information is indicative of the expected response of a patient to a method of treatment. Methods of determining relevant variance information and additional methods of using such variance information are also described.

10 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full	Title Oit:	ation Front	Review	Classification	Cate	Reference			Claims	[*)(d)()	Diranet E
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	L19 A1	ND inhala	tion the	rapy						8	

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Previous Page Next Page Go to Doc#

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Search Results - Record(s) 1 through 23 of 23 returned.

1. Document ID: US 20040185507 A1

Using default format because multiple data bases are involved.

L21: Entry 1 of 23

File: PGPB

Sep 23, 2004

PGPUB-DOCUMENT-NUMBER: 20040185507

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040185507 A1

TITLE: Anti-integrin antibodies, compositions, methods and uses

PUBLICATION-DATE: September 23, 2004

INVENTOR-INFORMATION:

COUNTRY RULE-47 STATE NAME CITY PAUS Giles-Komar, Jill Downingtown Snyder, Linda Pottstown PA US Trikha, Mohit Paoli PA US PA US Malvern Nakada, Marian T.

US-CL-CURRENT: 435/7.2; 530/388.22

Full Title Citat	ion Front F	Review Classification	Date	Reference	Sequences	Attachments	Claims	[6(4)]	ferane fe

2. Document ID: US 20040120952 A1

L21: Entry 2 of 23

File: PGPB

Jun 24, 2004

PGPUB-DOCUMENT-NUMBER: 20040120952

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040120952 A1

TITLE: Anti-TNF antibodies and peptides of human tumor necrosis factor

PUBLICATION-DATE: June 24, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Knight, David M. Berwin PA US Shealy, David J. Downingtown PA US

US-CL-CURRENT: 424/145.1

ABSTRACT:

Anti-TNF antibodies, fragments and regions thereof which are specific for human tumor necrosis factor-.alpha. (TNF.alpha.) and are useful in vivo diagnosis and therapy of a number of TNF.alpha.-mediated pathologies and conditions, as well as polynucleotides coding for murine and chimeric antibodies, methods of producing the antibody, methods of use of the anti-TNF antibody, or fragment, region or derivative thereof, in immunoassays and immunotherapeutic approaches are provided.

	Full	Titl∈	Citation	Frent	Review	Classification	Cate	Reference	Sequences	Attachments	Claime	15040	Draw, De
***************************************		2 1	Doguma	nt ID:	115 20	040077648	Λ 1		·				
			y 3 of		03 20	040077046	ΛI	File: P	GPB		Apr	22, 2	2004

PGPUB-DOCUMENT-NUMBER: 20040077648

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040077648 A1

TITLE: Methods and compositions of novel triazine compounds

PUBLICATION-DATE: April 22, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Timmer, Richard T.	Decatur	GA	US	
Alexander, Christopher W.	Norcross	GA	US	
Pillarisetti, Sivaram	Norcross	GA	US	
Saxena, Uday	Atlanta	GA	US	
Campbell, Karen A.	Durham	NC	US	

US-CL-CURRENT: 514/241; 544/212, 544/223

ABSTRACT:

The present invention relates to methods and compositions comprising compounds that treat pathophysiological conditions arising from inflammatory responses. In particular, the present invention is directed to compounds that inhibit or block glycated protein produced induction of the signaling-associated inflammatory response in endothelial cells. The present invention relates to compounds that inhibit smooth muscle proliferation. In particular, the present invention is directed to compounds that inhibit smooth muscle cell proliferation by modulating HSPGs such as Perlecan. The present invention further relates to the use of compounds to treat vascular occlusive conditions characterized by smooth muscle proliferation such as restenosis and atherosclerosis.

ull	Title	Citation	Front	Review	Classification	Irat∈	Reference	Sequences	Attachments:	Claime	1000	[-rand
IME		(Cathor)	Front	E & olelot	t lass means n	l'at≈	Reference	SECTION 100	Authornience	1.1311112	1 0000	Litan

4. Document ID: US 20040058412 A1

L21: Entry 4 of 23

File: PGPB

Mar 25, 2004

PGPUB-DOCUMENT-NUMBER: 20040058412

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040058412 A1

TITLE: Cell populations which co-express CD49c and CD90

PUBLICATION-DATE: March 25, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Ho, Tony W.	Berwyn	PA	US	
Kopen, Gene C.	Wynnewood	PA	US	
Righter, William F.	Ridley Park	PA	US	
Rutkowski, J. Lynn	Wynnewood	PA	US	
Wagner, Joseph	West Chester	PA	US	
Herring, W. Joseph	Valley Forge	PA	US	
Ragaglia, Vanessa	Newtown Square	PA	US	

US-CL-CURRENT: 435/69.1; 424/93.7, 435/320.1, 435/325, 435/366

ABSTRACT:

Substantially homogenous cells populations which co-express CD49c, CD90 and telomerase are made. In one embodiment, humans suffering from a degenerative, traumatic, acute injury, cardiac or neurological condition are treated with the substantially homogenous cells populations which co-express CD49c, CD90 and telomerase. In another embodiment, committed progenitor cells are made are made by selecting from a cultured source of a cell population which co-express CD49c and CD90 and modifying the cell population. The committed progenitor cells can be employed to treat a human suffering from a degenerative, traumatic, acute injury, cardiac or neurological condition and to formulate pharmaceutical compositions. In a further embodiment, a substantially homogenous populaton of cells which co-express CD49c, CD90 and at least one cardiac-related transcription factor is made and can be used to treat a human suffering from a cardiac condition.

Full	Titl∈	Citation	Frent	Review	Classification	Crate	Reference	Sequences	Attachments	Claims	150010	Draw, D.
П	5 D	ocume	nt ID:	US 20	030198954	A 1						

PGPUB-DOCUMENT-NUMBER: 20030198954

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030198954 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: October 23, 2003

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Bejanin, Stephane

Paris

FR

Tanaka, Hiroaki

Antony

FR

US-CL-CURRENT: 435/6; 536/23.2

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full	Titl≘	Citation	Front	Review	Massification	Cate	Referenc	e Sequences	Attachments	Claime	Finito	firant fir
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shirt da	6. I	Documen	t ID:	US 200	30181379	A1						
L21:	Entry	y 6 of 2	3				File:	PGPB		Sep	25,	2003

PGPUB-DOCUMENT-NUMBER: 20030181379

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030181379 A1

TITLE: Novel fibroblast growth factor (FGF23) and methods for use

PUBLICATION-DATE: September 25, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Econs, Michael	Indianapolis	IN	US	
White, Ken	Carmel	IN	US	
Strom, Tim Matthias	Munchen		DE	
Meitinger, Thomas	Munchen		DE	

US-CL-CURRENT: 514/12; 435/320.1, 435/325, 435/69.4, 530/399, 536/23.5

ABSTRACT:

The invention relates to novel nucleic acids encoding a fibroblast growth factor-23 (FGF23) and proteins encoded thereby, mutations in which are associated with autosomal dominant rickets (ADHR). The invention further relates to methods of diagnosing and treating hypophosphatemic and hyperphosphatemic disorders comprising inhibiting or stimulating, respectively, the biological activity of FGF23 in a patient. The invention also relates to methods of treating osteoporosis, dermatomyositis, and coronary artery disease comprising stimulating the biological activity of FGF23 in a patient.

Full	Titl≞	Citation	Front	Review	Classification	[rate	Reference	Sequences	Attachments	Claims	Pigit	Отзыс Он
	 7.	Docume	ent ID:	US 200	030176317	A1						
L21:	Entr	y 7 of	23				File: P	GPB		Sep	18,	2003

PGPUB-DOCUMENT-NUMBER: 20030176317

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030176317 A1

TITLE: Stabilization of hypoxia inducible factor (HIF) alpha

PUBLICATION-DATE: September 18, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Guenzler-Pukall, Volkmar	San Leandro	CA	US	
Neff, Thomas B.	Atherton	CA	US	
Wang, Qingjian	Davis	CA	US	
Arend, Michael P.	San Mateo	CA	US	
Flippin, Lee A.	Woodside	CA	US	
Melekhov, Alex	San Mateo	CA	US	

US-CL-CURRENT: 514/1

ABSTRACT:

The present invention relates to methods of stabilizing the alpha subunit of hypoxia inducible factor (HIF). The invention further relates to methods of preventing, pretreating, or treating conditions associated with HIF, including ischemic and hypoxic conditions. Compounds for use in these methods are also provided.

Full	Titl∈	Citation	Front Re	endem (Classification	Date	Freference	Sequences	Attachments	Claims	Pield	[traint [tr
	8.	Documen	nt ID: U	S 200	30170628	3 A1						
L21:	Entr	v 8 of 2	23				File: F	PGPB		Sep	11,	2003

PGPUB-DOCUMENT-NUMBER: 20030170628

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030170628 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: September 11, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bejanin, Stephane Tanaka, Hiroaki Paris Antony FR FR

ny

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/69.1, 435/7.1, 530/350, 530/388.1, 536/23.5

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full	Title	E Citation	Front	Review	Classification	() ata	Reference	Sequences	.4ttachments	Claima	Emic	[1350] [1-
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	9.	Docume	nt ID:	US 20	030162186	A 1						
L21:	Ent	ry 9 of	23				File:	PGPB		Aug	28,	2003

PGPUB-DOCUMENT-NUMBER: 20030162186

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030162186 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: August 28, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Bejanin, Stephane

Paris

FR

Tanaka, Hiroaki

Antony

FR

US-CL-CURRENT: 435/6; 435/183, 435/320.1, 435/325, 435/69.1, 536/23.2

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full	Titl≞	Citation	Front	Review	Classification	Cate Reference	Sequences	Ættachmenta	Claims	Dult	Отане
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L21: Entry 10 of 23

File: PGPB

Aug 21, 2003

PGPUB-DOCUMENT-NUMBER: 20030157485

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030157485 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: August 21, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bejanin, Stephane Paris FR Tanaka, Hiroaki Antony FR

US-CL-CURRENT: 435/6; 435/226, 435/320.1, 435/325, 435/69.1, 435/7.2, 530/388.26, 536/23.2, 800/8

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full T	Γitle	Citation	Front	Review	Classification	Date	Reference	Sequences	,4ttachmenta	Claims	Finis;	[trase [c
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☐ 11. Document ID: US 20030152921 A1

L21: Entry 11 of 23

File: PGPB

Aug 14, 2003

PGPUB-DOCUMENT-NUMBER: 20030152921

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030152921 A1

TITLE: Full-length human cDNAs encoding potentially secreted proteins

PUBLICATION-DATE: August 14, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Dumas Milne Edwards, Jean-Baptiste Paris FR
Bougueleret, Lydie Petit Lancy CH
Jobert, Severin Paris FR

US-CL-CURRENT: 435/6; 435/183, 536/23.2

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET

products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full Title Citation Front Flevieur Classification Date Flererence Sequences Attachments Claims Dobb Graw Co

12. Document ID: US 20030096247 A1

L21: Entry 12 of 23

File: PGPB

May 22, 2003

PGPUB-DOCUMENT-NUMBER: 20030096247

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030096247 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: May 22, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bejanin, Stephane Paris FR Tanaka, Hiroaki Antony FR

US-CL-CURRENT: 435/6; 435/183, 435/320.1, 435/325, 435/69.1, 530/350, 536/23.2, 800/8

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims 1990	fit3hd
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☐ 13. Document ID: US 20030092011 A1

L21: Entry 13 of 23

File: PGPB

May 15, 2003

PGPUB-DOCUMENT-NUMBER: 20030092011

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030092011 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: May 15, 2003

INVENTOR-INFORMATION:

COUNTRY RULE-47 CITY STATE NAME

FR Bejanin, Stephane Paris FR Tanaka, Hiroaki Antony

US-CL-CURRENT: 435/6; 435/183, 435/320.1, 435/325, 435/69.1, 435/7.9, 536/23.2,

800/3

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full	Title Citation Fro	ont Fleviese Classification	Date Reference	Sequences	Attachments	Claims	Elide)ť	[1] [1]
	14. Document	ID: US 2003007273	7 A1					
L21:	Entry 14 of 23	3	File:	PGPB		Apr :	17,	2003

PGPUB-DOCUMENT-NUMBER: 20030072737

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030072737 A1

TITLE: Tissue protective cytokines for the protection, restoration, and enhancement of responsive cells, tissues and organs

PUBLICATION-DATE: April 17, 2003

INVENTOR-INFORMATION:

STATE COUNTRY RULE-47 CITY NAME Brines, Michael Woodbridge CTUS Croton On Hudson NY US Cerami, Antony Cerami, Carla Sleepy Hollow NY US

US-CL-CURRENT: 424/85.1; 530/351

ABSTRACT:

Methods and compositions are provided for protecting or enhancing a responsive cell, tissue, organ or body part function or viability in vivo, in situ or ex vivo in mammals, including human beings, by systemic or local administration of a tissue protective cytokine.

Full	Titl⊕	Citation	Front	Review	Classification	Date Reference	Sequences	#.ttachments	Claima	F100 ()	[arana [a
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Page 10 of 15

☐ 15. Document ID: US 20030040044 A1

L21: Entry 15 of 23

File: PGPB

Feb 27, 2003

PGPUB-DOCUMENT-NUMBER: 20030040044

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030040044 A1

TITLE: Anti-dual integrin antibodies, compositions, methods and uses

PUBLICATION-DATE: February 27, 2003

INVENTOR - INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Heavner, George	Malvern	PA	US	
Giles-Komar, Jill	Downingtown	PA	US	
Snyder, Linda	Pottstown	PA	US	
Trikha, Mohit	Paoli	PA	US	

US-CL-CURRENT: 435/69.1; 424/146.1, 435/320.1, 435/326, 530/387.2, 530/388.26, 536/23.53

ABSTRACT:

The present invention relates to at least one novel anti-dual integrin antibodies, including isolated nucleic acids that encode at least one anti-dual integrin antibody, dual integrin, vectors, host cells, transgenic animals or plants, and methods of making and using thereof, including therapeutic compositions, methods and devices.

Full	Title	Citation	Front	Review	Classification	Cate	Reference	Sequences	Attachmenta	Claims	Ejodí;	Drawe De
	16	Docume	ent ID	. IIC 2	.003002724	8 Δ1	manifican opensyllinastyly distr	•		<u></u>		· · · · · · · · · · · · · · · · · · ·
		y 16 of		. 052	.005002721	0111	File:	PGPB		Feb	6,	2003

PGPUB-DOCUMENT-NUMBER: 20030027248

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030027248 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: February 6, 2003

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47
Bejanin, Stephane Paris FR
Tanaka, Hiroaki Antony FR

US-CL-CURRENT: 435/69.1; 435/183, 435/320.1, 435/325, 435/6, 530/350, 536/23.2

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full Title	CHESTION	Front	Review	Classification	[rate	Reference	Sequences	Attachmenta	Claims	Detic	Orano E
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17. Document ID: US 20030027161 A1

L21: Entry 17 of 23

File: PGPB

Feb 6, 2003

PGPUB-DOCUMENT-NUMBER: 20030027161

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030027161 A1

TITLE: Human cDNAs and proteins and uses thereof

PUBLICATION-DATE: February 6, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bejanin, Stephane Paris FR Tanaka, Hiroaki Antony FR

US-CL-CURRENT: 435/6; 435/183, 435/320.1, 435/325, 435/69.1, 530/350, 536/23.2, 800/8

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full	Titl∈	Citation	Front	Regiena	Classification	Late	Reference	Sequences	Attachments	Claims	[104]	Draw, D
	18.	Docume	ent ID	: US 2	2003002221	0 A 1						
L21:	Entr	y 18 of	23				File:	PGPB		Jan	30,	2003

PGPUB-DOCUMENT-NUMBER: 20030022210

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030022210 A1

Record List Display

TITLE: T cell induced tissue repair and regeneration

PUBLICATION-DATE: January 30, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bonyhadi, Mark Issaquah WA US Berenson, Ron Mercer Island WA US

US-CL-CURRENT: 435/6; 424/93.7, 435/368

ABSTRACT:

The present invention relates to methods for the use of T cells or supernatants therefrom, and more particularly, activated T cells, in facilitating and/or regulating the differentiation, de-differentiation, maturation, organization, repair, and regeneration of various cells/tissues. Methods for inducing tissue repair and regeneration in vitro and in vivo are disclosed. The present invention also relates to compositions of cells, including activated T cells and/or cells resulting from the co-culture with activated T cells, and their use in inducing tissue repair and regeneration in vivo.

19. Document ID: US 20020165191 A1

L21: Entry 19 of 23 File: PGPB Nov 7, 2002

PGPUB-DOCUMENT-NUMBER: 20020165191

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020165191 A1

TITLE: Spatial and temporal control of gene expression using a heat shock protein promoter in combination with local heat

PUBLICATION-DATE: November 7, 2002

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Moonen, Chrit Bordeaux FR

US-CL-CURRENT: 514/44; 607/108

ABSTRACT:

The invention provides methods for using local heat to control gene expression. The heat shock protein (hsp) gene promoter is recombined with a selected therapeutic gene and expressed in selected cells. Local controlled heating is used to activate the hsp promoter, for example by using focused ultrasound controlled by MRI.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	#.ttachinento	Claims	DetC	Drame De
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	20.	Docum	ent ID	: US 2	002015600	1 A1	•					
L21:	Entr	y 20 of	23				File:	PGPB		Oct	24,	2002

PGPUB-DOCUMENT-NUMBER: 20020156001

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020156001 A1

TITLE: Novel fibroblast growth factor (FGF23) and methods for use

PUBLICATION-DATE: October 24, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Econs, Michael	Indianapolis	IN	US	
White, Ken	Carmel	IN	US	
Strom, Tim Matthias	Munchen		DE	
Meitinger, Thomas	Munchen		DE	

US-CL-CURRENT: 514/12; 435/320.1, 435/325, 435/6, 435/69.1, 530/399, 536/23.5

ABSTRACT:

The invention relates to novel nucleic acids encoding a fibroblast growth factor-23 (FGF23) and proteins encoded thereby, mutations in which are associated with autosomal dominant rickets (ADHR). The invention further relates to methods of diagnosing and treating hypophosphatemic and hyperphosphatemic disorders comprising inhibiting or stimulating, respectively, the biological activity of FGF23 in a patient. The invention also relates to methods of treating osteoporosis, dermatomyositis, and coronary artery disease comprising stimulating the biological activity of FGF23 in a patient.

Full	Titl∈	Citation	Front	Review	©lassification	() ate	Reference	Sequences	.4ttachments	Claims	1,3000	Crawt Co-

	21.	Docume	ent ID	: US 2	002010260	4 A1						
L21:	Entr	y 21 of	23				File:	PGPB		Aug	1,	2002

PGPUB-DOCUMENT-NUMBER: 20020102604

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020102604 A1

TITLE: Full-length human cDNAs encoding potentially secreted proteins

PUBLICATION-DATE: August 1, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Milne Edwards, Jean-Baptiste Dumas

Paris

FR

Bougueleret, Lydie Jobert, Severin

Petit Lancy Paris

CH FR

US-CL-CURRENT: <u>435/7.1</u>; 530/350, 536/23.1

ABSTRACT:

The invention concerns GENSET polynucleotides and polypeptides. Such GENSET products may be used as reagents in forensic analyses, as chromosome markers, as tissue/cell/organelle-specific markers, in the production of expression vectors. In addition, they may be used in screening and diagnosis assays for abnormal GENSET expression and/or biological activity and for screening compounds that may be used in the treatment of GENSET-related disorders.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims 1990 Draw, D.

22. Document ID: US 6794363 B2

L21: Entry 22 of 23

File: USPT

Sep 21, 2004

COUNTRY

US-PAT-NO: 6794363

DOCUMENT-IDENTIFIER: US 6794363 B2

TITLE: Isolated amyloid inhibitor protein (APIP) and compositions thereof

DATE-ISSUED: September 21, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE

Bejanin; Stephane Paris FR Tanaka; Hiroaki Antony FR

US-CL-CURRENT: 514/12; 435/23, 530/350, 536/23.5

ABSTRACT:

The invention provides polynucleotides and polypeptides encoding an isolated amyloid inhibitor protein (APIP) and compositions thereof. The polypeptides of the subject invention can be used to inhibit the catabolism or sequential cleavage of arnyloid beta precursor protein (APP) by Sequential cleavage of APP by beta secretase and gamma secretase.

10 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title Citation Front Review Classification Cate Reference of Experience Olaina 1500 Grane [o 23. Document ID: US 6239172 B1

L21: Entry 23 of 23

File: USPT

May 29, 2001

US-PAT-NO: 6239172

DOCUMENT-IDENTIFIER: US 6239172 B1

** See image for Certificate of Correction **

TITLE: Formulations for treating disease and methods of using same

DATE-ISSUED: May 29, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Kaesemeyer; Wayne H.

Augusta

. GA

US-CL-CURRENT: 514/460

ABSTRACT:

A therapeutic mixture comprised of L-arginine and angiogenic growth factors is disclosed for the treatment of diseases related to endothelial dysfunction.

14 Claims, 1 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 1

Full	Title Citation From	l Review	Classification	[/ate	Reference	A STATE OF THE STA		Claims	[564];	Drawe D
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Search Results - Record(s) 1 through 7 of 7 returned.

1. Document ID: US 20040185507 A1

Using default format because multiple data bases are involved.

L22: Entry 1 of 7

File: PGPB

Sep 23, 2004

Jun 24, 2004

PGPUB-DOCUMENT-NUMBER: 20040185507

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040185507 A1

TITLE: Anti-integrin antibodies, compositions, methods and uses

PUBLICATION-DATE: September 23, 2004

INVENTOR-INFORMATION:

RULE-47 CITY STATE COUNTRY NAME PA US Downingtown Giles-Komar, Jill US Snyder, Linda Pottstown PΑ Trikha, Mohit Paoli PA US PA US Nakada, Marian T. Malvern

US-CL-CURRENT: 435/7.2; 530/388.22

Full	Title	Citation	Front	Fleview	Claszinication	Date	Reference	Sequence:	Attachments	Claims	1,3640	[Habet [H
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File: PGPB

PGPUB-DOCUMENT-NUMBER: 20040120952

PGPUB-FILING-TYPE: new

L22: Entry 2 of 7

DOCUMENT-IDENTIFIER: US 20040120952 A1

TITLE: Anti-TNF antibodies and peptides of human tumor necrosis factor

PUBLICATION-DATE: June 24, 2004

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47
Knight, David M. Berwin PA US

Shealy, David J. Downingtown PA US

US-CL-CURRENT: 424/145.1

ABSTRACT:

Anti-TNF antibodies, fragments and regions thereof which are specific for human tumor necrosis factor-.alpha. (TNF.alpha.) and are useful in vivo diagnosis and therapy of a number of TNF.alpha.-mediated pathologies and conditions, as well as polynucleotides coding for murine and chimeric antibodies, methods of producing the antibody, methods of use of the anti-TNF antibody, or fragment, region or derivative thereof, in immunoassays and immunotherapeutic approaches are provided.

Full Title Citation Front Review Class	itication Date Reference Sequences 3	Attachments (Claims 1500) Draw De
☐ 3. Document ID: US 200400	77648 A1	
L22: Entry 3 of 7	File: PGPB	Apr 22, 2004

PGPUB-DOCUMENT-NUMBER: 20040077648

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040077648 A1

TITLE: Methods and compositions of novel triazine compounds

PUBLICATION-DATE: April 22, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Timmer, Richard T.	Decatur	GA	US	
Alexander, Christopher W.	Norcross	GA	US	
Pillarisetti, Sivaram	Norcross	GA	us	
Saxena, Uday	Atlanta	GA	US	
Campbell, Karen A.	Durham	NC	US	

US-CL-CURRENT: 514/241; 544/212, 544/223

ABSTRACT:

The present invention relates to methods and compositions comprising compounds that treat pathophysiological conditions arising from inflammatory responses. In particular, the present invention is directed to compounds that inhibit or block glycated protein produced induction of the signaling-associated inflammatory response in endothelial cells. The present invention relates to compounds that inhibit smooth muscle proliferation. In particular, the present invention is directed to compounds that inhibit smooth muscle cell proliferation by modulating HSPGs such as Perlecan. The present invention further relates to the use of compounds to treat vascular occlusive conditions characterized by smooth muscle proliferation such as restenosis and atherosclerosis.

Full Title 0	Citation Front	Review	Olazzifikation	(ate	Reference	Sequences	#Hachments	Ctaime	[56b)	Lirano, Di

4. Document ID: US 20030096754 A1

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L22: Entry 4 of 7

File: PGPB

May 22, 2003

PGPUB-DOCUMENT-NUMBER: 20030096754

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030096754 A1

TITLE: Vascular endothelial growth factor variants

PUBLICATION-DATE: May 22, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Pollitt, N. Stephen Los Altos CA US Abraham, Judith A. San Jose CA US

US-CL-CURRENT: 514/12

ABSTRACT:

The invention is directed to a method of enhancing the biological activity of vascular endothelial growth factors ($\underline{\text{VEGF}}$). The invention further concerns certain $\underline{\text{VEGF}}$ variants having enhanced biological activity, methods and means for preparing these variants, and pharmaceutical compositions comprising them. In a further aspect, the invention concerns methods of treatment using, and articles of manufacture containing such VEGF variants.

Full	Title Citation Front	Review Classification	Date Refere	ancel Sequences	#ttachments	Claima	15640	[41300] [4
	5. Document ID:	US 20030040044	A1					
L22:	Entry 5 of 7		File:	PGPB		Feb	27,	2003

PGPUB-DOCUMENT-NUMBER: 20030040044

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030040044 A1

TITLE: Anti-dual integrin antibodies, compositions, methods and uses

PUBLICATION-DATE: February 27, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Heavner, George	Malvern	PA	US	
Giles-Komar, Jill	Downingtown	PA	US	
Snyder, Linda	Pottstown	PA	US	
Trikha, Mohit	Paoli	PA	US	

US-CL-CURRENT: 435/69.1; 424/146.1, 435/320.1, 435/326, 530/387.2, 530/388.26, 536/23.53

ABSTRACT:

Record List Display Page 4 of 5

The present invention relates to at least one novel anti-dual integrin antibodies, including isolated nucleic acids that encode at least one anti-dual integrin antibody, dual integrin, vectors, host cells, transgenic animals or plants, and methods of making and using thereof, including therapeutic compositions, methods and devices.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims 1990	f.17300 f.

6. Document ID: US 20020019350 A1

L22: Entry 6 of 7

File: PGPB

Feb 14, 2002

PGPUB-DOCUMENT-NUMBER: 20020019350

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020019350 A1

TITLE: Targeted angiogenesis

PUBLICATION-DATE: February 14, 2002

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RUL	E-47
Levine, Arnold J. New York NY US	
Mitterer, Artur Orth, Donau AT	
Falkner, Falko-Guenter Orth, Donau AT	
Scheiflinger, Friedrich Vienna AT	
Dorner, Friedrich Vienna AT	

US-CL-CURRENT: 514/12; 530/399

ABSTRACT:

The invention relates to compositions, methods, and gene therapy reagents to promote or to inhibit angiogenesis in the treatment of peripheral vascular or cardiovascular diseases, utilizing a chimeric molecule comprising an angiogenic factor linked to a targeting molecule that specifically binds to a vascular endothelium.

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☐ 7. Document ID: US 6475796 B1

L22: Entry 7 of 7

File: USPT

Nov 5, 2002

US-PAT-NO: 6475796

DOCUMENT-IDENTIFIER: US 6475796 B1

TITLE: Vascular endothelial growth factor variants

DATE-ISSUED: November 5, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Pollitt; N. Stephen

Los Altos

CA

Abraham; Judith A.

San Jose

CA

US-CL-CURRENT: 435/455; 424/198.1, 514/2, 530/350

ABSTRACT:

The invention is directed to a method of enhancing the biological activity of vascular endothelial growth factors (VEGF). The invention further concerns certain VEGF variants having enhanced biological activity, methods and means for preparing these variants, and pharmaceutical compositions comprising them. In a further aspect, the invention concerns methods of treatment using, and articles of manufacture containing such VEGF variants.

17 Claims, 17 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 17

Full	Title Citation	Front	Review	Classification	Frate	Reference	Maria and American Services		Claima	15000	[113m)
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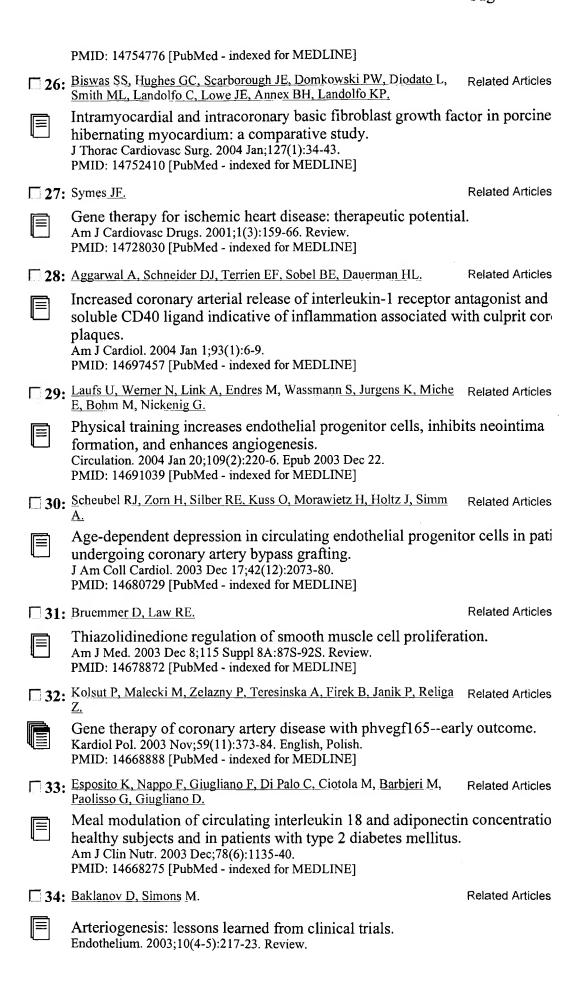


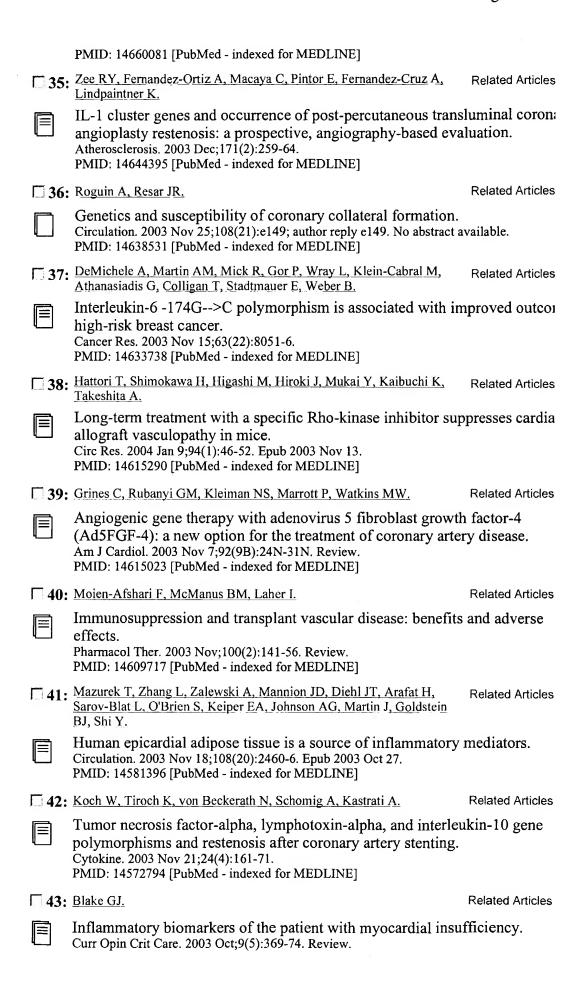


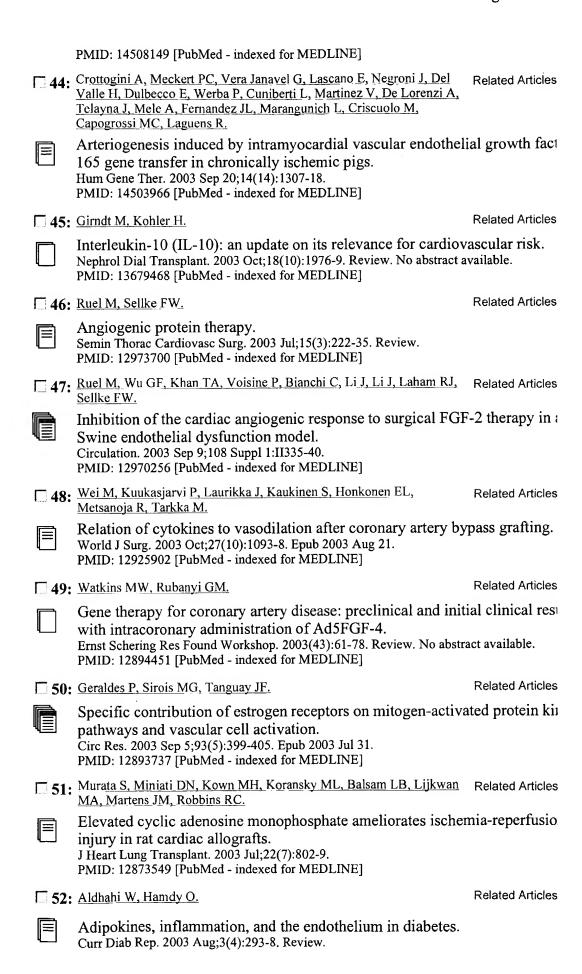
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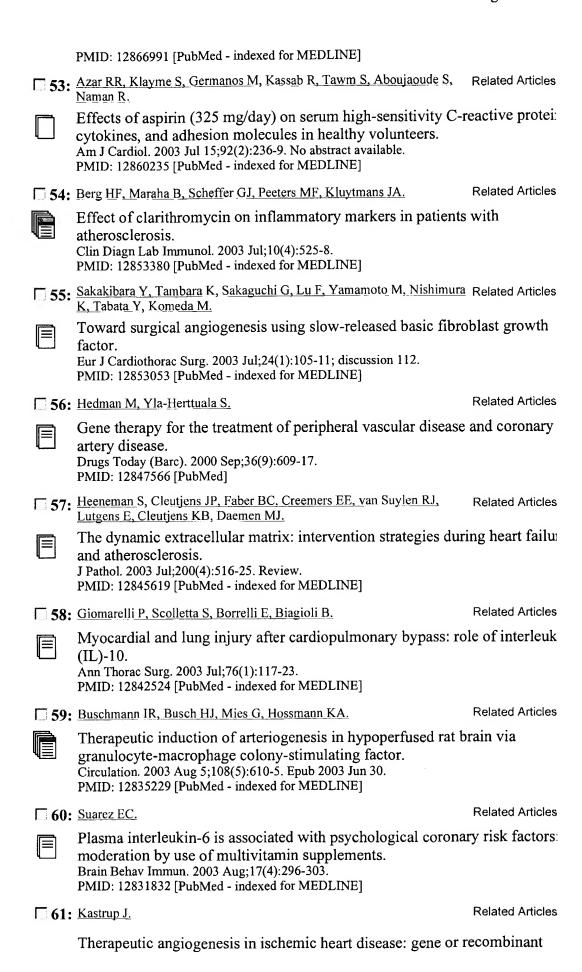
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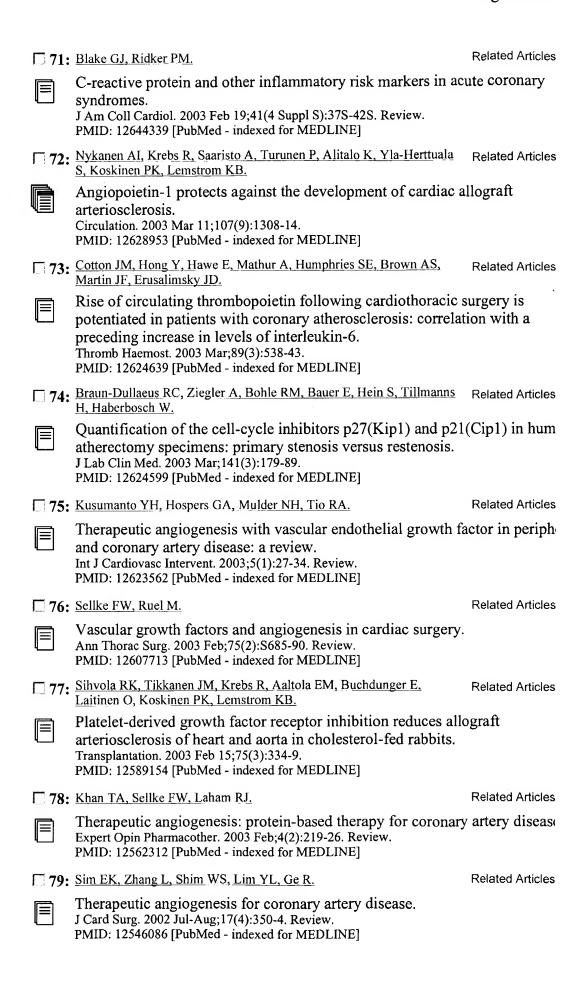








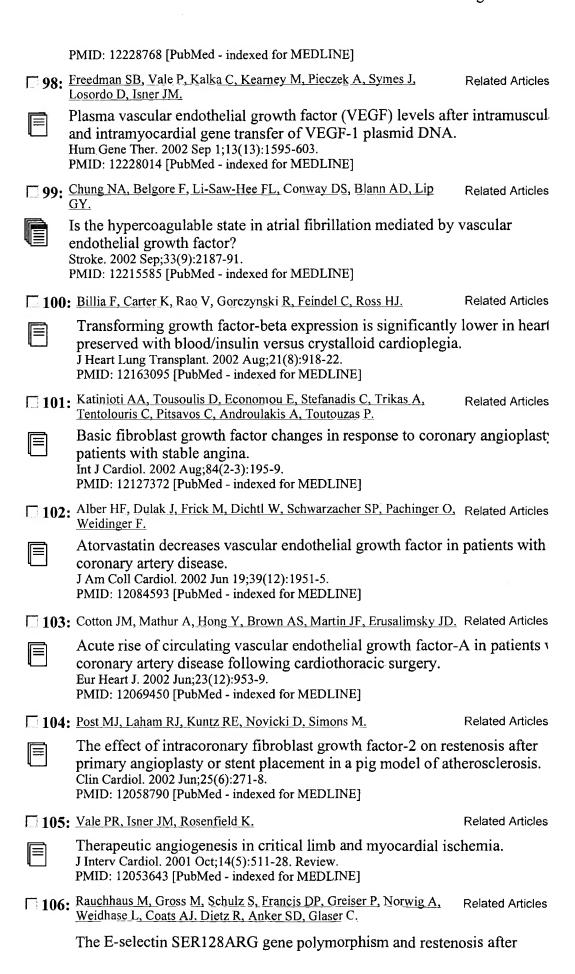
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	Therapeutic angiogenesis: review of current concepts and fut J Heart Lung Transplant. 2003 Apr;22(4):370-82. Review. No abstract av PMID: 12681415 [PubMed - indexed for MEDLINE]	ure directions. ailable.
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	Translational physiology: porcine models of human coronary implications for preclinical trials of therapeutic angiogenesis. J Appl Physiol. 2003 May;94(5):1689-701. Review. PMID: 12679343 [PubMed - indexed for MEDLINE]	artery disease
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	Serum level of the antiinflammatory cytokine interleukin-10 prognostic determinant in patients with acute coronary syndro Circulation. 2003 Apr 29;107(16):2109-14. Epub 2003 Mar 31. PMID: 12668510 [PubMed - indexed for MEDLINE]	is an importan omes.



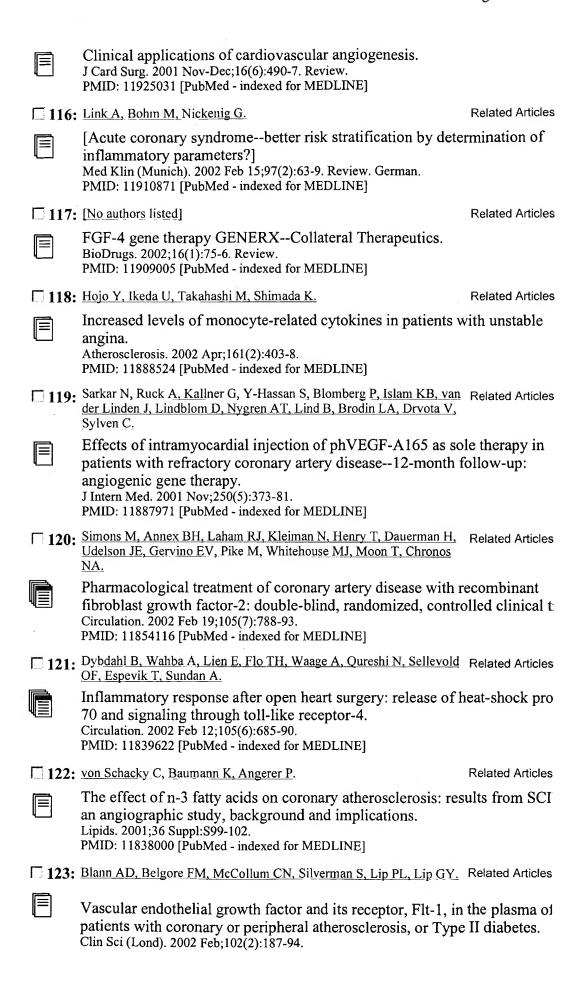
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□ 82:	<u>Leinonen E, Hurt-Camejo E, Wiklund O, Hulten LM, Hiukka A, Taskinen MR.</u>	Related Articles
	Insulin resistance and adiposity correlate with acute-phase reacell adhesion molecules in type 2 diabetes. Atherosclerosis. 2003 Feb;166(2):387-94. PMID: 12535753 [PubMed - indexed for MEDLINE]	action and solu
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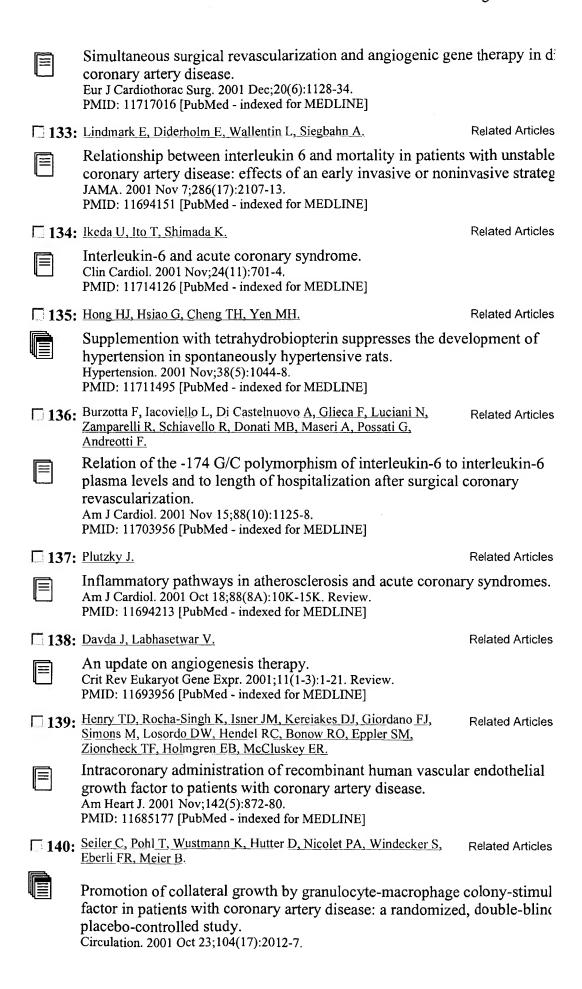
□ 89:	Rasmussen HS, Rasmussen CS, Macko J, Yonehiro G.	Related Articles
	Angiogenic gene therapy strategies for the treatment of cardic Curr Opin Mol Ther. 2002 Oct;4(5):476-81. Review. PMID: 12435049 [PubMed - indexed for MEDLINE]	ovascular dise
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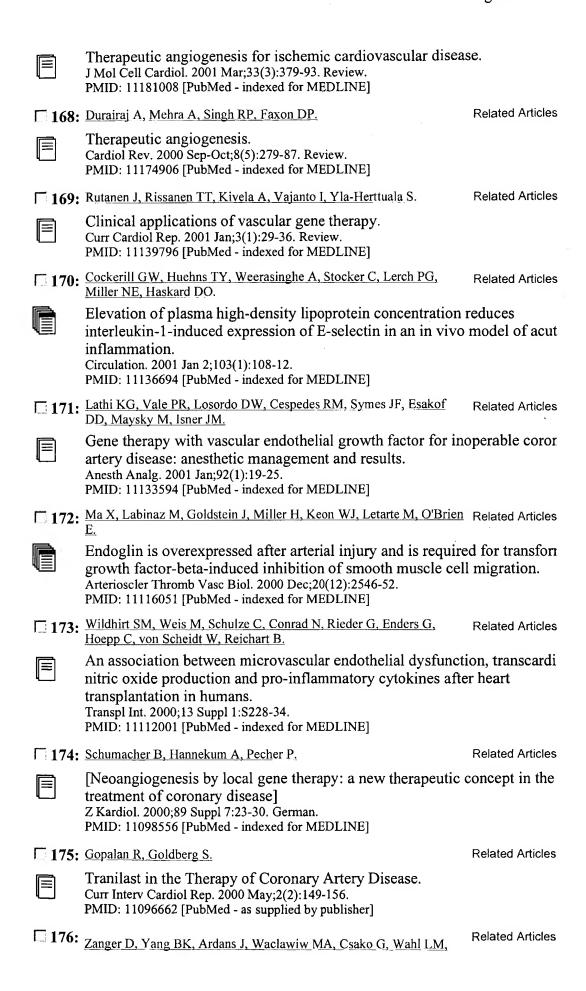
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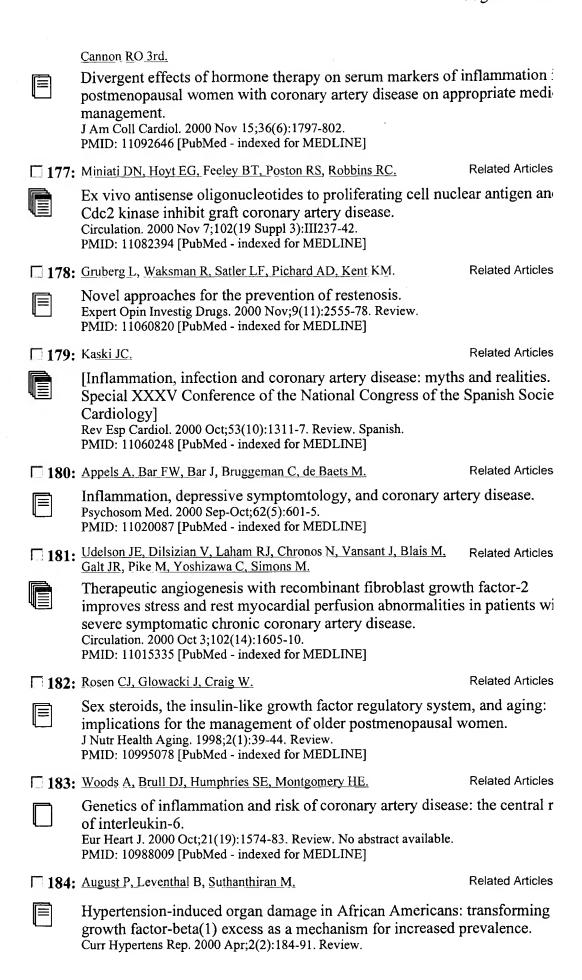


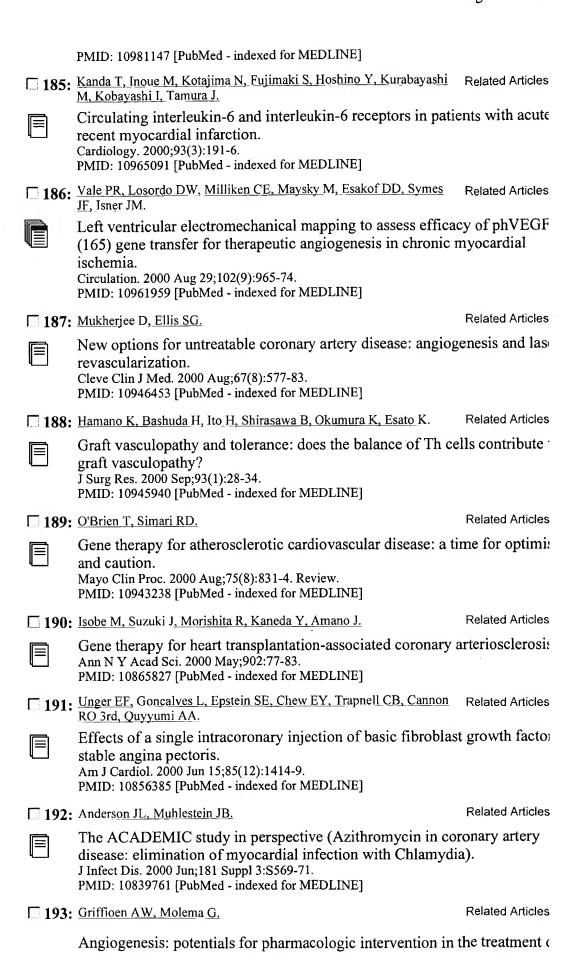
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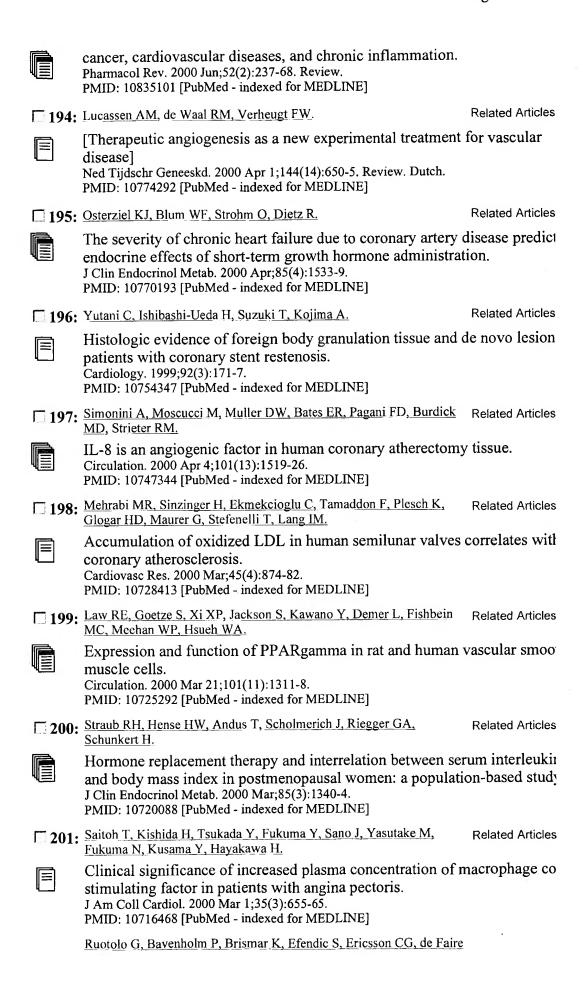
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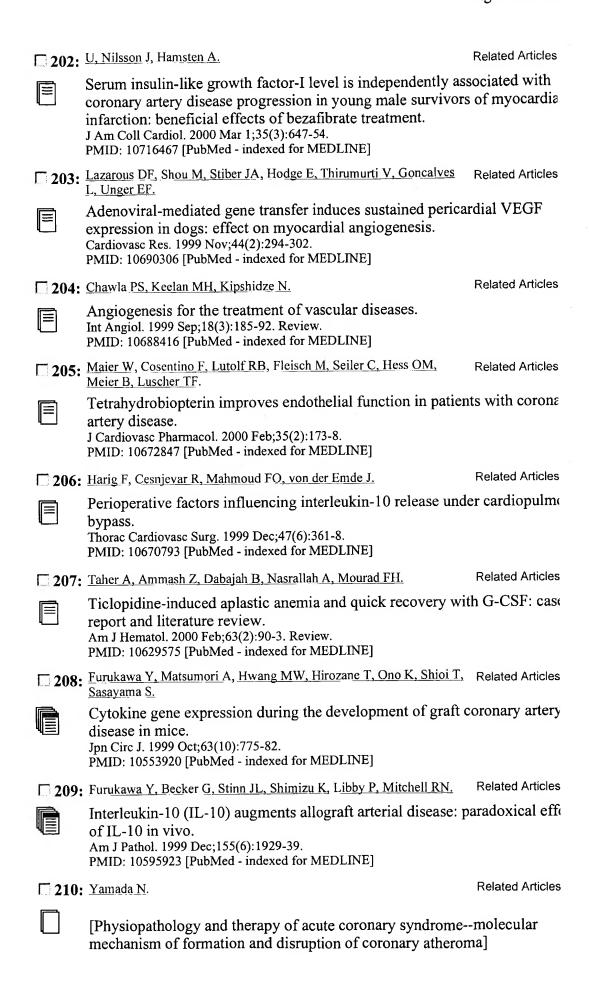
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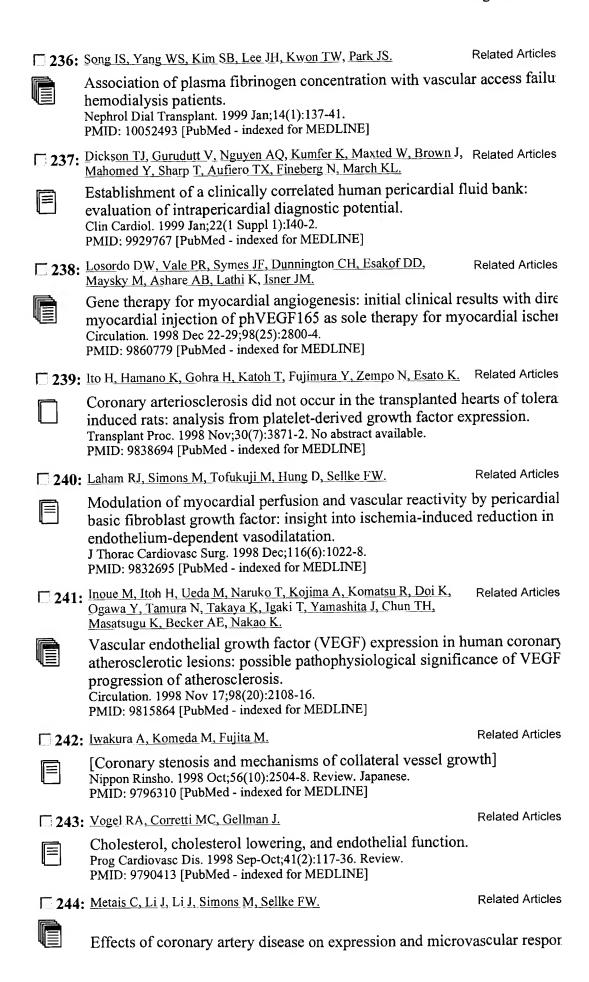
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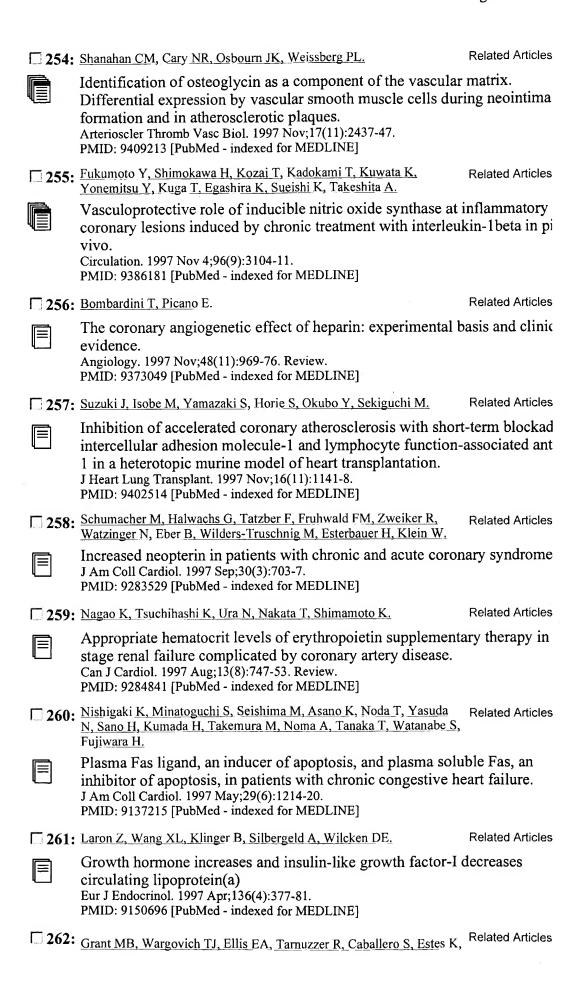
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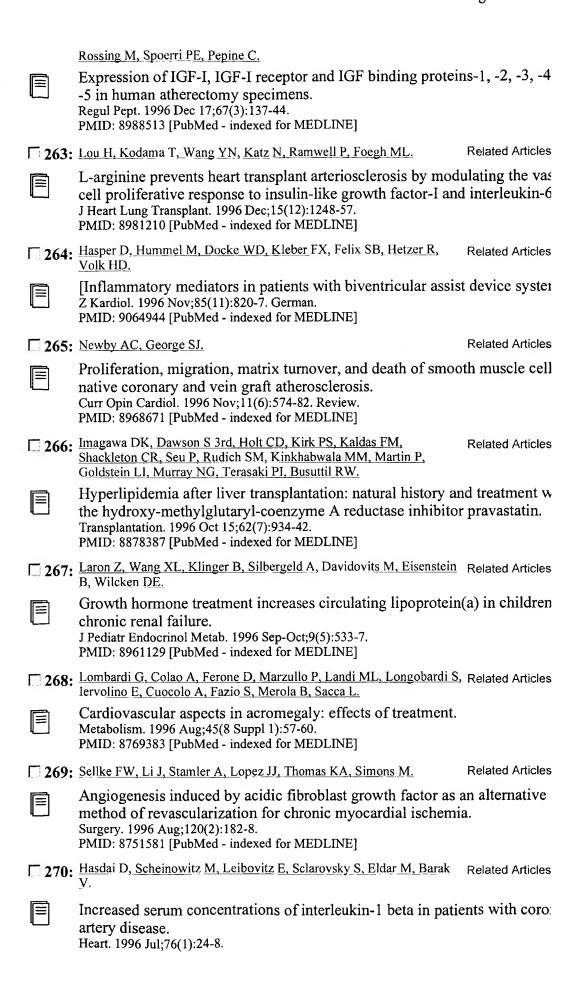
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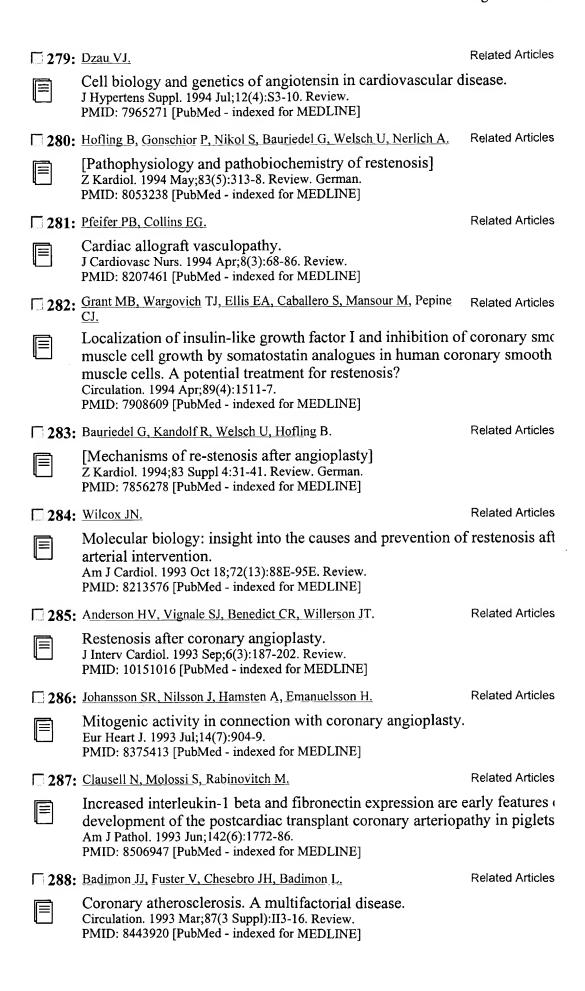


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Nora R, Abrams JS, Tait NS, Hiponia DJ, Silverman HJ.

Department of Medicine, University of Maryland Cancer Center, Baltimore 2

Arterial and pulmonary artery catheters were used to monitor the cardiopulm effects of recombinant interleukin-2 (rIL-2) given iv at a dose of 100,000 U/l every 8 hours on days 1-5 to 10 patients with metastatic solid tumors. As anticipated, a severe capillary leak syndrome developed in all patients. Myoc infarction (MI) occurred unexpectedly in three patients, as evidenced by a foinjury pattern on ECG and elevations of creatinine phosphokinase myocardia band fractions. All patients receiving rIL-2 exhibited major reductions in their ventricular stroke work index (47 +/- 11 g.m/m2 to 29 +/- g.m/m2), an index cardiac contractility. It remains uncertain whether the MIs were a byproduct capillary leak syndrome in patients with underlying coronary artery disease c whether rIL-2 directly or indirectly damages cardiac muscle.

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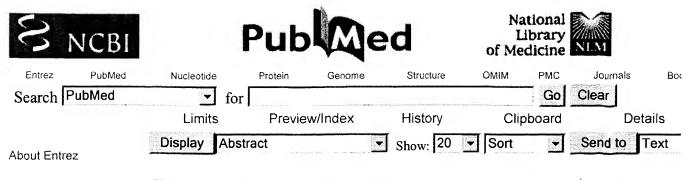
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Therapeutic angiogenesis with basic fibroblast growth factor: technique and early results.

Sellke FW, Laham RJ, Edelman ER, Pearlman JD, Simons M.

Angiogenesis Research Center, Department of Surgery at Beth Israel Deacon Medical Center, Boston, Massachusetts 02215, USA. fsellke@bidmc.harvarc

BACKGROUND: Patients not amenable to complete myocardial revascularia by conventional methods present a difficult clinical problem. Here we presen early results and technical considerations of the administration of basic fibrol growth factor for the induction of collateral growth using heparin-alginate sle release devices in patients undergoing coronary artery bypass grafting. METHODS: Eight patients were enrolled. Patients were candidates if they have least one graftable obstructed coronary artery and at least one major arterial distribution not amenable to revascularization, a serum creatinine level less the mg/dL, ejection fraction greater than 0.20, and estimated operative mortality less than 25%. During conventional coronary artery bypass grafting, 10 hepa alginate devices, each containing either 1 microg or 10 microg of basic fibrol growth factor, were implanted in the epicardial fat in multiple regions of the unrevascularizable territory and also in the distal distribution of a grafted or I artery. RESULTS: There was no mortality and no evidence of renal, hematol or hepatic toxicity during follow-up. Three months after the operation, all par remain free of angina. Seven patients were examined with stress perfusion sc Three patients had clear enhancement of perfusion to the unrevascularized myocardium, 1 patient had a new fixed defect, and 3 had minimal overall cha but had evidence of new small, fixed perfusion defects. Seven patients had improved or similar myocardial contractile function (ejection fraction at 3-m follow-up = 0.53 + - 0.22 versus 0.47 + - 0.14 preoperatively). One patient suffered a perioperative myocardial infarction in the area of basic fibroblast growth factor administration. CONCLUSIONS: This preliminary study demonstrates the safety and technical feasibility of therapeutic angiogenesis basic fibroblast growth factor delivered by heparin-alginate slow-release dev Further studies examining the safety, clinical efficacy, and long-term results ongoing.

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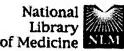
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Nucleotide OMIM **PMC** Protein Structure Journals Books Search PubMed for langina AND CK-MB Go Clear Limits Preview/Index History Clipboard Details Display Summary Show: 500 × Sort Send to Text About Entrez Items 1 - 304 of 304 One t **Text Version** 1: Sousa JM, Hermann JL, Guimaraes JB, Menezes PP, Carvalho AC. Related Articles, L Evaluation of systolic, diastolic, and pulse pressure as risk factors for severe Entrez PubMed coronary arteriosclerotic disease in women with unstable angina non-ST-Overview Help | FAQ elevation acute myocardial infarction. Tutorial Arq Bras Cardiol. 2004 May;82(5):430-3, 426-9. Epub 2004 Jun 08. English, Portuguese. New/Noteworthy PMID: 15340673 [PubMed - in process] E-Utilities 2: Frossard M, Fuchs I, Leitner JM, Hsieh K, Vlcek M, Losert H, Related Articles, L Domanovits H, Schreiber W, Laggner AN, Jilma B. **PubMed Services** Journals Database Platelet function predicts myocardial damage in patients with acute myocardia MeSH Database infarction. Single Citation Matcher **Batch Citation Matcher** Circulation. 2004 Sep 14;110(11):1392-7. Epub 2004 Aug 16. PMID: 15313953 [PubMed - in process] Clinical Queries LinkOut 3: Gaspardone A, De Fabritiis P, Scaffa R, Nardi P, Palombi F, Versaci F, Cubby Related Articles, L Chiariello L. Related Resources [Stem cell mobilization after coronary artery bypass grafting] Order Documents Ital Heart J. 2004 Jan;5(1 Suppl):23-8. Italian. **NLM Catalog** PMID: 15253141 [PubMed - in process] **NLM Gateway TOXNET** 4: Wang YN, Che SM, Ma AQ. Related Articles, L Consumer Health Clinical Alerts Clinical significance of serum cytokines IL-1beta, sIL-2R, IL-6, TNF-alpha, a ClinicalTrials.gov IFN-v in acute coronary syndrome. PubMed Central Chin Med Sci J. 2004 Jun; 19(2): 120-4. PMID: 15250248 [PubMed - indexed for MEDLINE] 5. Canos DA, Mintz GS, Berzingi CO, Apple S, Kotani J, Pichard AD, Satler Related Articles, L LF, Suddath WO, Waksman R, Lindsay J Jr, Weissman NJ. Clinical, angiographic, and intravascular ultrasound characteristics of early saphenous vein graft failure. J Am Coll Cardiol. 2004 Jul 7;44(1):53-6. PMID: 15234406 [PubMed - indexed for MEDLINE] 1 6: Uva MS, Rodrigues V, Monteiro N, Pereira F, Bervens D, Caria R, Related Articles, L. Mesquita A, Pedro A, Bau J, Matias F, Magalhaes MP. Coronary surgery: which method to use? Rev Port Cardiol. 2004 Apr;23(4):517-30. English, Portuguese. PMID: 15224641 [PubMed - in process] 7: Arslanagic A, Gerc V. Related Articles, L [New approach in the diagnosis of acute coronary syndrome] Med Arh. 2004;58(2 Suppl 1):43-5. Review. Bosnian. PMID: 15202307 [PubMed - indexed for MEDLINE] 8: Sardella G, De Luca L, Adorisio R, Di Russo C, Fedele F. Related Articles. L Effects of rotational atherectomy with a reduced burr-to-artery ratio on corona no-reflow.

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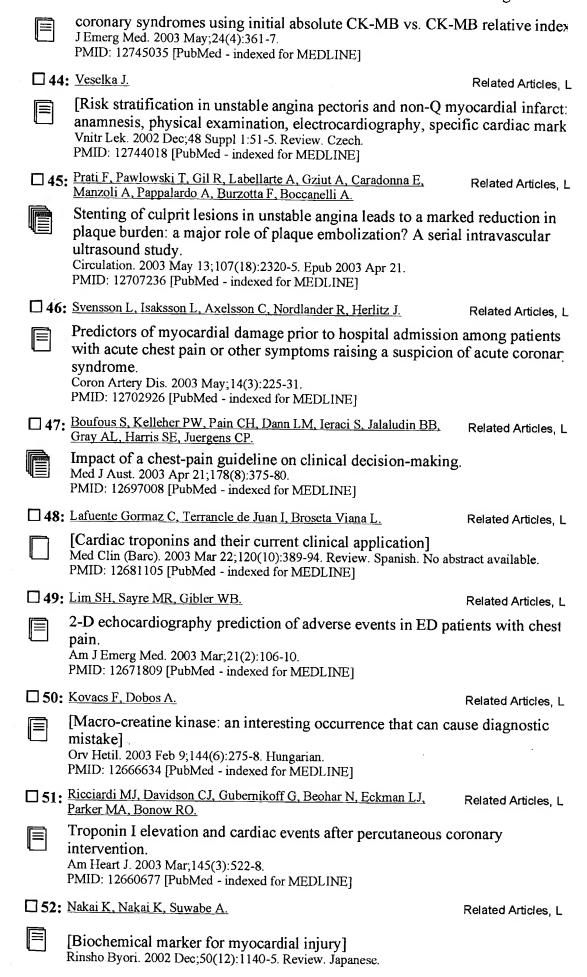
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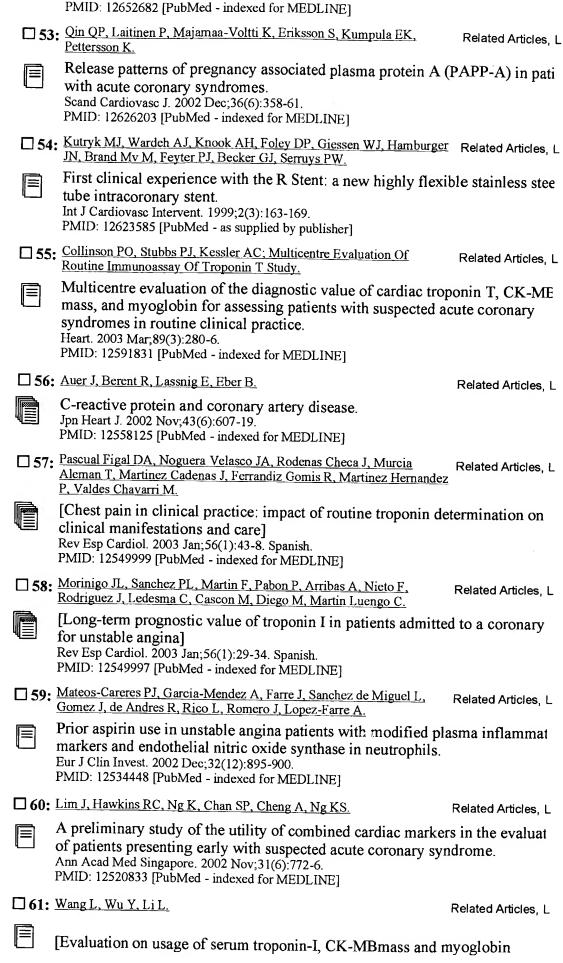
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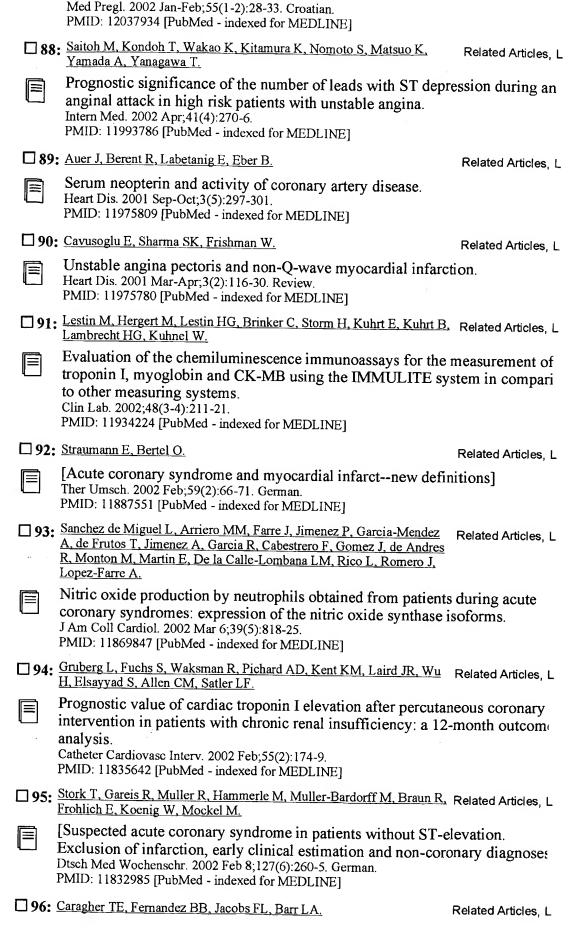


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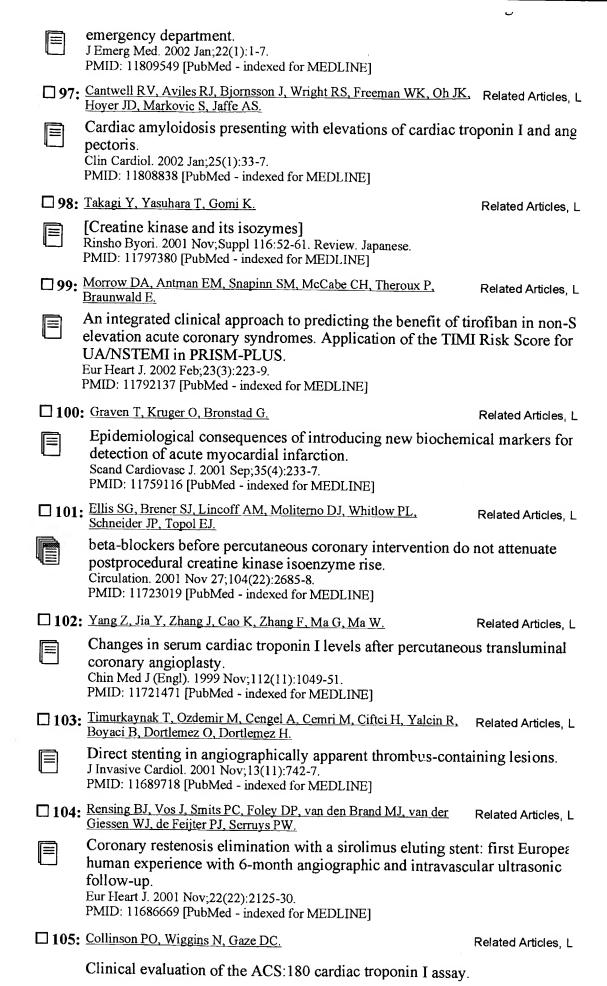
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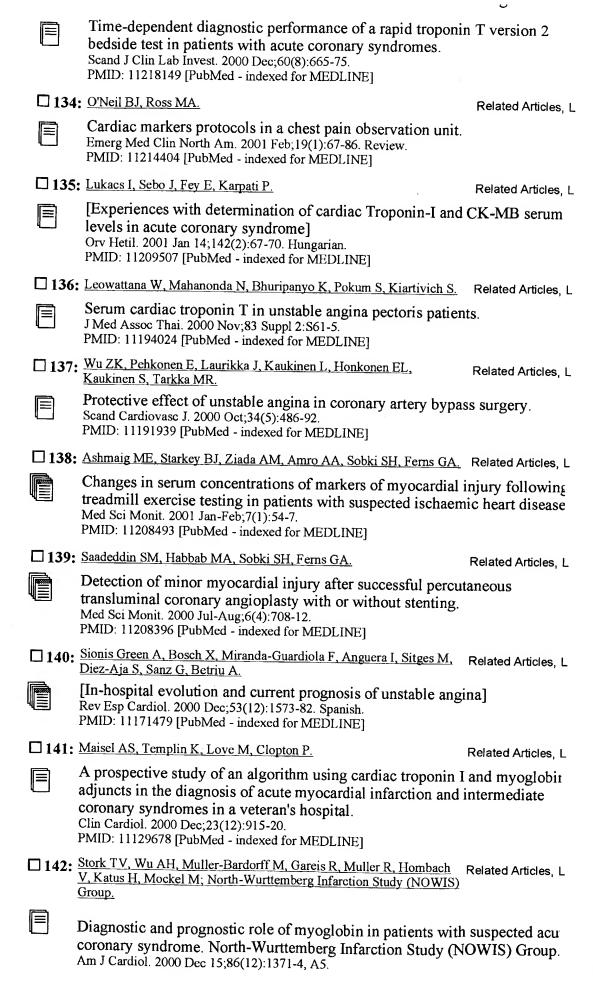
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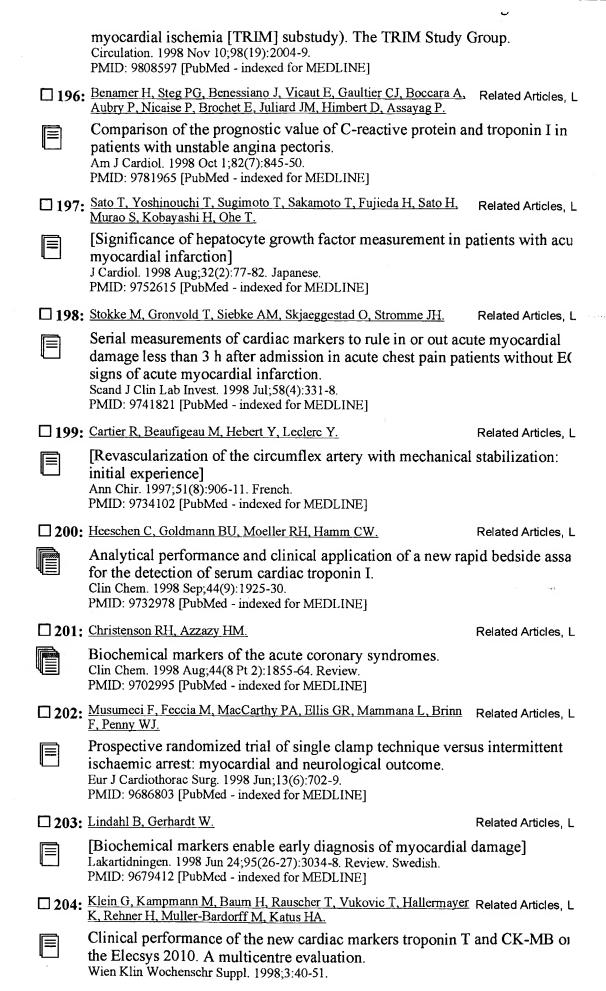
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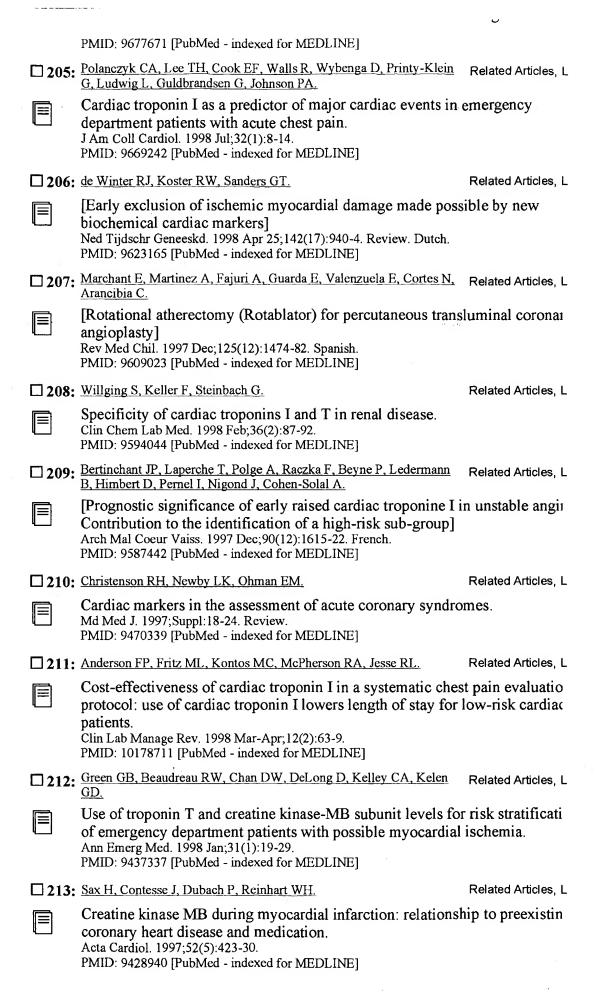
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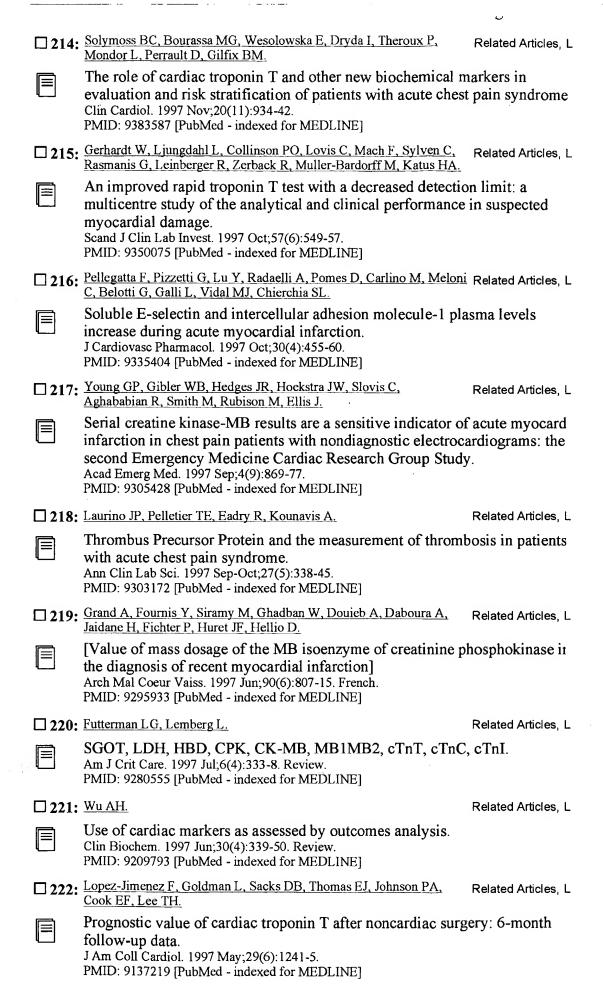
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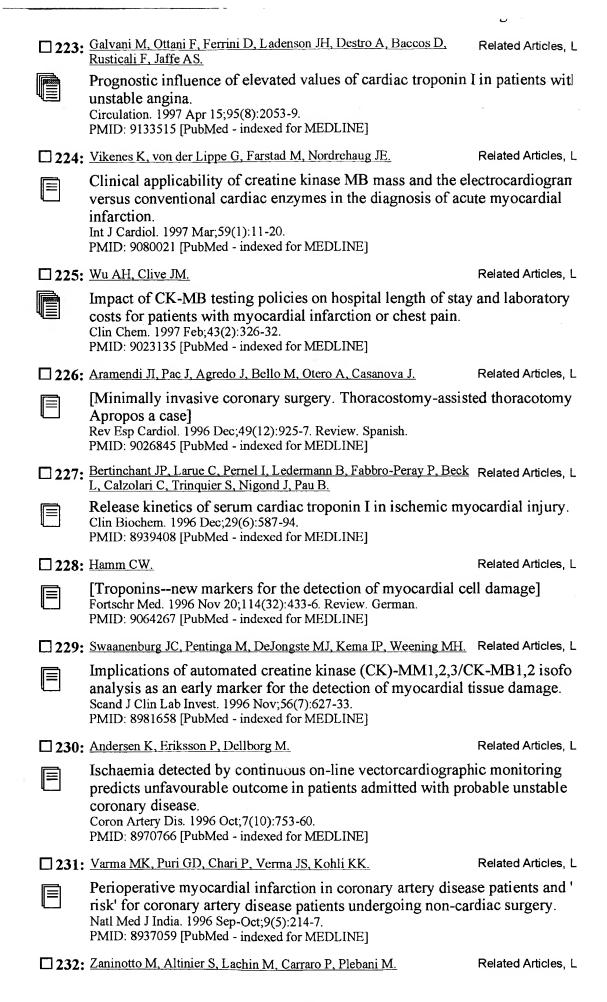
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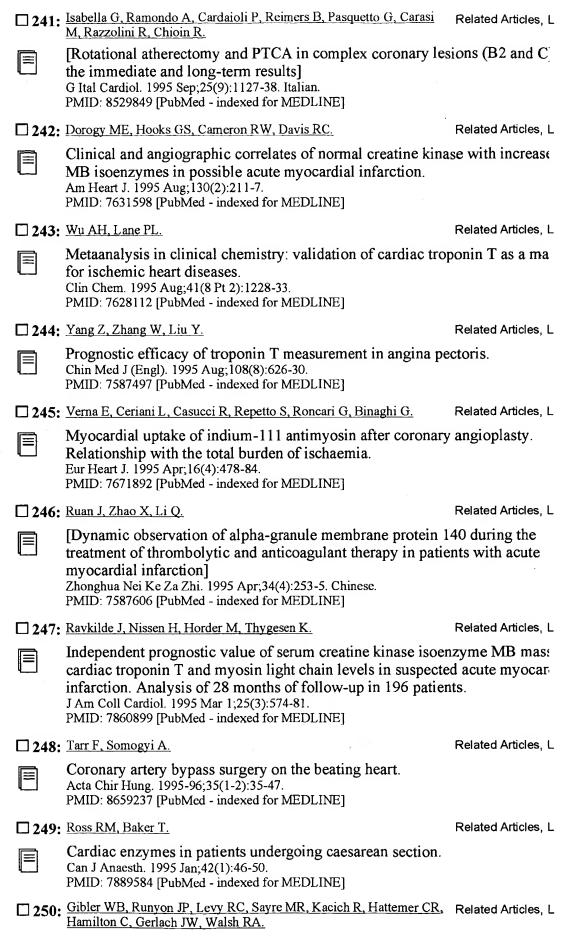




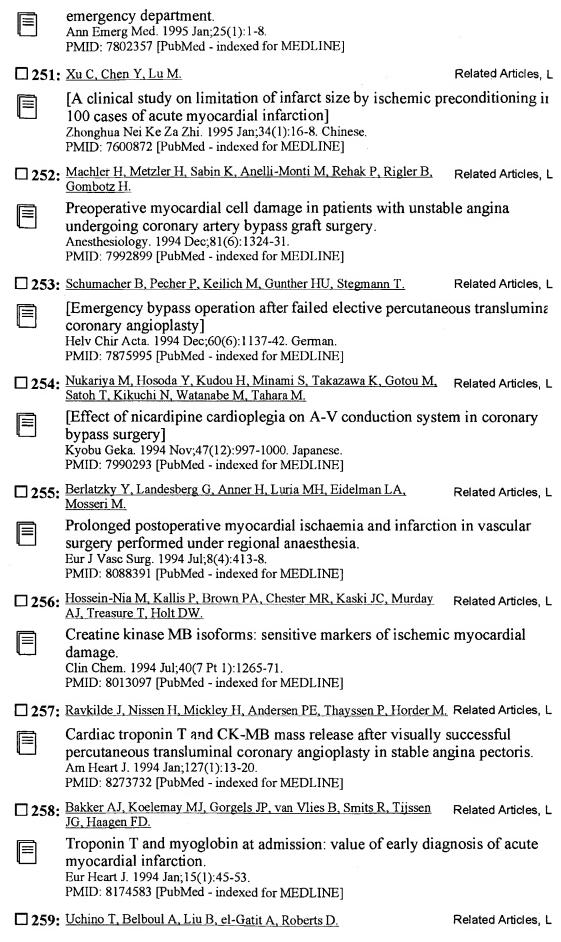


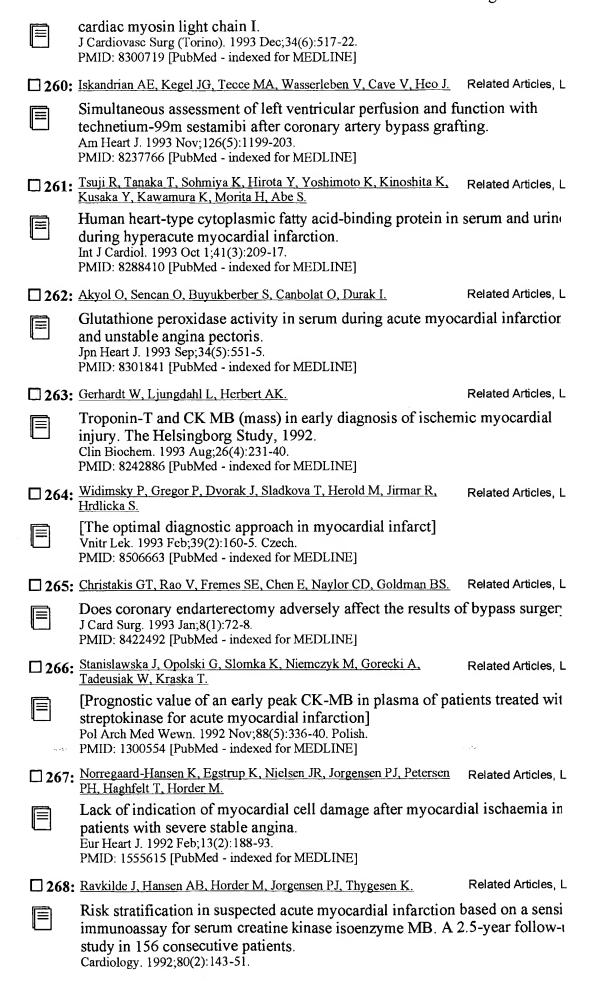


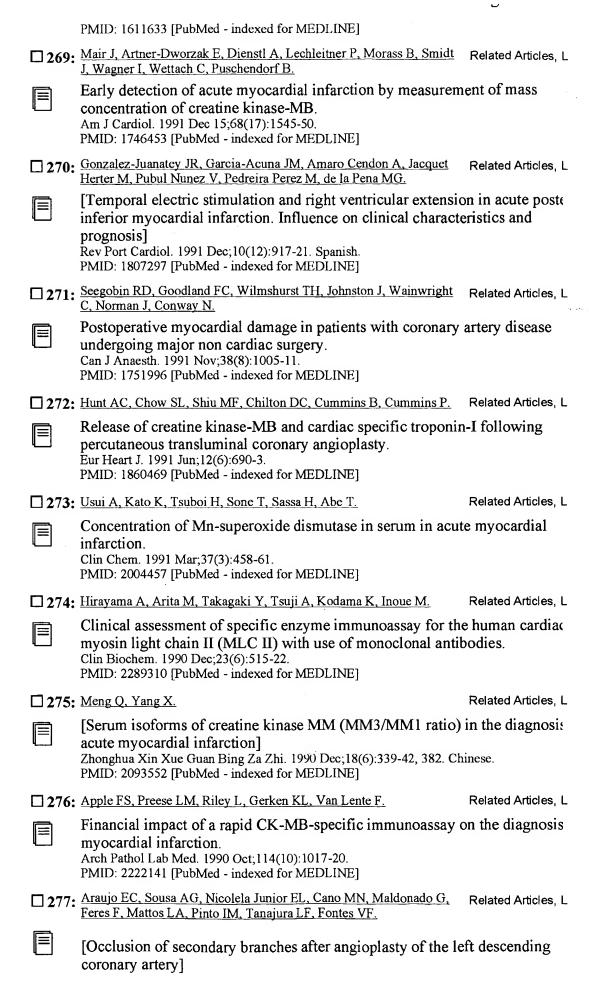
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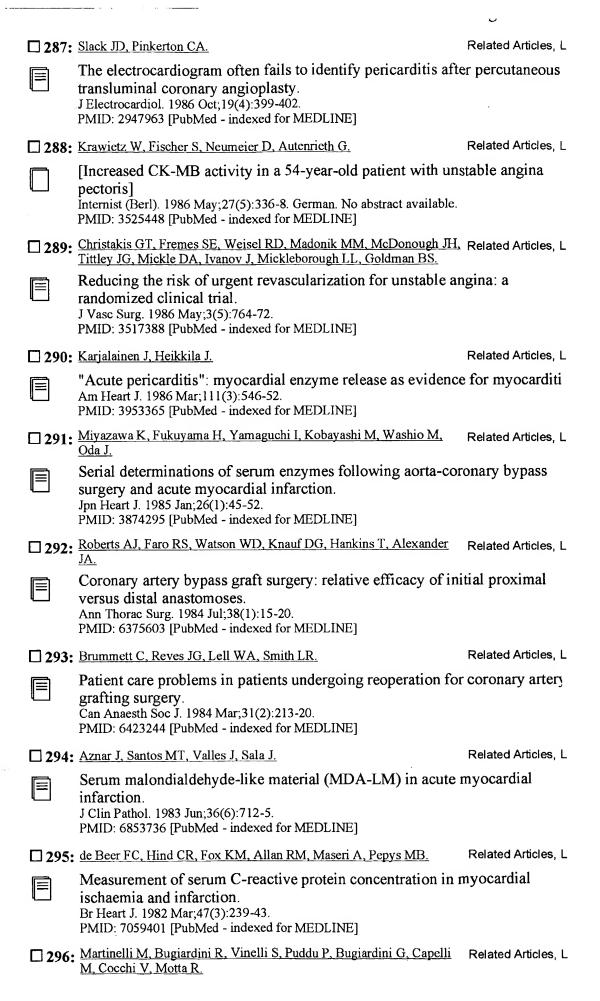
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                     Gene Delivery System In Two Phase II Gene Therapy
Angioplasty Clinical Trials
                     BioAccess, ( ***1 Apr 1998*** ) pp. N/A.
SOURCE:
                     ISSN: 1356-3432.
                     English
LANGUAGE:
WORD COUNT:
                        712
                     *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
L7
     ANSWER 10 OF 78 PROMT COPYRIGHT 2004 Gale Group on STN
                     1998:173353 PROMT
ACCESSION NUMBER:
                     Genentech Starts Trials Of Recombinant
                                                                ***VEGF***
TITLE:
                                      ***13 Apr 1998*** ) pp. N/A.
SOURCE:
                     Marketletter,
                     ISSN: 0951-3175.
                     English
LANGUAGE:
                        756
WORD COUNT:
                     *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
L7
     ANSWER 11 OF 78 PROMT COPYRIGHT 2004 Gale Group on STN
                     1998:143219 PROMT
ACCESSION NUMBER:
TITLE:
                     GENEMEDICINE Proprietary Cationic Lipid Gene Delivery
                     System Is Employed in Two Phase II Gene Therapy Angioplasty
                     clinical Trials.
                                        ***19 Mar 1998*** ) pp. 3190068.
SOURCE:
                     Business Wire, (
                     English
LANGUAGE:
WORD COUNT:
                     *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
     ANSWER 12 OF 78 PROMT COPYRIGHT 2004 Gale Group on STN
L7
ACCESSION NUMBER:
                     1998:41293 PROMT
                     Genentech's Year-End Results Show Growth Plan on Track:
TITLE:
                     Earnings Increase Nine Percent on Revenues Exceeding $1
                     Billion.
                     Business Wire, ( ***22 Jan 1998*** ) pp. 01220134.
SOURCE:
                     English
LANGUAGE:
                       2029
WORD COUNT:
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*FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
     ANSWER 13 OF 78 PROMT COPYRIGHT 2004 Gale Group on STN
L7
                        1998:41215
                                       PROMT
ACCESSION NUMBER:
                        ZENECA AIMS TO DOUBLE R & D OUTPUT BY 2002
TITLE:
                        Pharmaceutical Business News, ( ***19 Dec 1997*** ) pp.
SOURCE:
                        ISSN: 0956-0661.
LANGUAGE:
                        English
                           1561
WORD COUNT:
                         *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
      ANSWER 14 OF 78 PROMT COPYRIGHT 2004 Gale Group on STN
L7
                         97:629445 PROMT
ACCESSION NUMBER:
                         Zeneca_Allays Fears Of Near-Term Product Gap
TITLE:
                                              ***8 Dec 1997*** ) pp. N/A.
                         Marketletter, (
SOURCE:
                         ISSN: 0951-3175.
                         English
LANGUAGE:
                           Ž214
WORD COUNT:
                         *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
      ANSWER 15 OF 78 PROMT COPYRIGHT 2004 Gale Group on STN
L7
                         97:97804 PROMT
ACCESSION NUMBER:
                         Biotech's Bellwethers
TITLE:
                         BioVenture View, ( ***1 Feb 1997*** ) pp. N/A.
SOURCE:
                         ISSN: 0892-1903.
                         English
LANGUAGE:
                           4435
WORD COUNT:
                         *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
      ANSWER 16 OF 78 USPATFULL on STN
L7
AN
         2001:33286 USPATFULL
         Prevention and treatment of cardiovascular pathologies with tamoxifen
TI
         analogues
         Grainger, David J., Cambridge, United Kingdom
IN
         Metcalfe, James C., Cambridge, United Kingdom
         Kunz, Lawrence L., Redmond, WA, United States
         Schroff, Robert W., Edmonds, WA, United States
NeoRx Corporation, Seattle, WA, United States (U.S. corporation)
US 6197789
B1 20010306
         US 6197789
PΙ
                        19961219
         wo 9640098
         us 1997-973570
                                       19971205 (8)
ΑI
         wo 1996-US10211
                                       19960607
                                                   PCT 371 date
                                       19980908
                                       19980908
                                                  PCT 102(e) date
         Continuation-in-part of Ser. No. US 1995-478936, filed on 7 Jun 1995, now abandoned Continuation-in-part of Ser. No. US 1995-476735, filed on 7 Jun 1995, now patented, Pat. No. US 5595722 Continuation-in-part of Ser. No. US 1995-477393, filed on 7 Jun 1995 Continuation-in-part of Ser. No. US 1995-486334, filed on 7 Jun 1995, now patented, Pat. No. US
RLI
         5770609
DT
         Utility
FS
         Granted
LN.CNT 4577
INCL
         INCLM: 514/319.000
         INCLS: 514/324.000; 514/422.000; 514/428.000; 514/444.000; 514/448.000;
                  514/651.000; 514/866.000
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514/324.000; 514/422.000; 514/428.000; 514/444.000; 514/448.000; 514/651.000; 514/866.000

<--

514/319; 514/324; 514/422; 514/428; 514/444; 514/448; 514/651; 514/866

Methods for the treatment and diagnosis of cardiovascular disease

20001205

Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.

514/319.000

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 17 OF 78 USPATFULL ON STN

2000:164259 USPATFULL

ICS: A61K031-40; A61K031-38; A61K031-135

Falb, Dean, Wellesley, MA, United States

ICM: A61K031-445

corporation)

us 6156500

NCL

IC

EXF

L7

AN

TI

IN

PA

ΡI

NCLM:

NCLS:

[7]

```
19950210 (8)
        us 1995-386844
ΑI
        Utility
DT
        Granted
LN.CNT 4817
        INCLM: 435/006.000
INCL
        INCLS: 436/501.000; 935/077.000
                435/006.000
NCL
        NCLM:
                436/501.000
        NCLS:
         [7]
IC
        ICM: C12Q001-68
        435/6; 435/810; 435/69.1; 435/7.1; 436/501; 436/63; 530/300; 530/350;
EXF
         530/387.1: 536/23.1: 536/24.1: 536/24.3-24.33: 935/77.78
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 18 OF 78 USPATFULL ON STN
L7
         2000:145865 USPATFULL
ΑN
         Targeted contrast agents for diagnostic and therapeutic use
TI
        Unger, Evan C., Tucson, AZ, United States
TN
        Fritz, Thomas A., Tucson, AZ, United States
Gertz, Edward W., Paradise Valley, AZ, United States
ImaRx Pharmaceutical Corp., Tucson, AZ, United States (U.S. corporation)
US 6139819 20001031 <--
PA
ΡI
         US 1997-932273
                                      19970917 (8)
ΑI
        Continuation-in-part of Ser. No. US 1996-660032, filed on 6 Jun 1996, now abandoned which is a continuation-in-part of Ser. No. US
RLI
         1996-640464, filed on 1 May 1996, now abandoned which is a
         continuation-in-part of Ser. No. US 1995-497684, filed on 7 Jun 1995, now abandoned And a continuation-in-part of Ser. No. US 1996-666129,
         filed on 19 Jun 1996, now patented, Pat. No. US 6033645
DT
         Utility
         Granted
FS
         7523
LN.CNT
         INCLM: 424/009.520
INCL
         INCLS: 424/009.510; 424/450.000
                 424/009.520
NCL
         NCLM:
         NCLS:
                 424/009.510; 424/450.000
         [7]
IC
         ICM: A61B008-00
         ICS: A61K009-127
         424/9.52; 424/9.51; 424/9.5; 424/450; 424/812; 600/441; 600/458; 264/4.1; 427/2.14; 427/213.3; 428/402.2
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 19 OF 78 USPATFULL ON STN
L7
         2000:138360 USPATFULL
ΑN
         Hydroxyl-containing bicyclic compounds
ΤI
         Underiner, Gail E., Brier, WA, United States
IN
         Porubek, David, Seattle, WA, United States
         Klein, J. Peter, Vashon Island, WA, United States
         Woodson, Paul, Edmonds, WA, United States
         Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
                                      20001017
PΙ
         us 6133274
                                       19961126 (8)
         us 1996-756703
ΑI
         Continuation of Ser. No. US 1993-153256, filed on 16 Nov 1993, now abandoned which is a continuation-in-part of Ser. No. US 1992-976353,
RLI
         filed on 16 Nov 1992, now patented, Pat. No. US 5473070
DT
         Utility
         Granted
FS
LN.CNT 1646
INCL
         INCLM: 514/263.000
         INCLS: 544/267.000
         NCLM:
                  514/263.360
NCL
                 544/267.000
         NCLS:
IC
         [7]
         ICM: C07D473-04
         ICS: A61K031-52
EXF
         544/267; 514/263
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 20 OF 78 USPATFULL ON STN
L7
         2000:128465 USPATFULL
AN
         Compositions and methods for treatment and diagnosis of cardiovascular
TI
         Falb, Dean A., Wellesley, MA, United States
IN
         Gimbrone, Jr., Michael Á., Jámaica Plain, MA, United States
Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
 PA
```

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corporation)
       Brigham and Women's Hospital, Boston, MA, United States (U.S.
       corporation)
PΙ
       US 6124433
                                 20000926
       US 1997-944496
                                 19971006 (8)
ΑI
       Division of Ser. No. US 1996-599654, filed on 9 Feb 1996, now patented,
RLI
       Pat. No. US 5882925 which is a continuation-in-part of Ser. No. US
       1995-485573, filed on 7 Jun 1995 which is a continuation-in-part of Ser.
       No. US 1995-386844, filed on 10 Feb 1995
       Utility
DT
       Granted
FS
LN.CNT 5924
       INCLM: 530/350.000
INCL
       INCLS: 530/324.000; 530/326.000; 536/023.100; 536/023.500; 435/069.100;
               435/320.100; 435/325.000
               530/350.000
NCL
       NCLM:
               435/069.100; 435/320.100; 435/325.000; 530/324.000; 530/326.000;
       NCLS:
               536/023.100; 536/023.500
       [7]
TC
       ICM: C07K016-00
       ICS: C12N015-00
       536/23.1; 536/24.1; 536/24.3; 536/23.5; 435/6; 435/69.1; 435/7.1; 435/172.3; 435/320.1; 435/325; 935/32; 935/52; 530/350
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 21 OF 78 USPATFULL on STN
ΑN
       2000:121539 USPATFULL
       Methods for regulating transcription factors
TI
       Qabar, Maher N., Redmond, WA, United States
IN
       McMillan, Michael K., Bellevue, WA, United States
       Kahn, Michael S., Kirkland, WA, United States
       Tulinsky, John E., Seattle, WA, United States
       Ogbu, Cyprian O., Bellevue, WA, United States
       Mathew, Jessymol, Bellevue, WA, United States
       Molecumetics Ltd., Bellevue, WA, United States (U.S. corporation)
PA
       US 6117896
                                  20000912
PΙ
ΑI
       US 1998-22934
                                 19980212 (9)
       Continuation-in-part of Ser. No. US 1997-797915, filed on 10 Feb 1997,
RLI
       now abandoned And a continuation-in-part of Ser. No. US 692420
       US 1997-47067P
                             19970519 (60)
PRAI
       Utility
DT
       Granted
FS
LN.CNT 4501
INCL
        INCLM: 514/384.000
        INCLS: 514/248.000; 530/323.000; 530/332.000; 548/263.400
               514/384.000
NCL
       NCLM:
               514/248.000; 530/323.000; 530/332.000; 548/263.400
       NCLS:
IC
        [7]
        ICM: A61K031-41
        ICS: C07K005-00; C07K007-00; C07K016-00; C07D249-12 514/248; 514/384; 530/332; 530/323; 548/263.4
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 22 OF 78 USPATFULL on STN
        2000:109372 USPATFULL
ΑN
ΤI
        In vivo agents comprising cationic drugs, peptides and metal chelators
        with acidic saccharides and glycosaminoglycans, giving improved
        site-selective localization, uptake mechanism, sensitivity and
       kinetic-spatial profiles, including tumor sites
IN
        Ranney, David F., Dallas, TX, United States
        Access Pharmaceuticals, Inc., Dallas, TX, United States (U.S.
PA
        corporation)
                                                                          <--
PI
                                  20000822
        us 6106866
        us 1995-509338
                                  19950731 (8)
ΑI
        Utility
DT
        Granted
FS
LN.CNT
        3913
        INCLM: 424/499.000
INCL
        INCLS: 424/489.000; 424/491.000; 424/493.000; 424/548.000; 514/054.000;
               514/062.000; 530/322.000; 536/054.000
               424/499.000
NCL
        NCLM:
               424/489.000; 424/491.000; 424/493.000; 424/548.000; 514/054.000; 514/062.000; 530/322.000; 536/054.000
        NCLS:
IC
        ICM: A61K031-726
        530/322; 424/548; 424/489; 424/491; 424/493; 424/499; 536/54; 514/54;
EXF
```

```
514/62
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 23 OF 78 USPATFULL ON STN
L7
        2000:105913 USPATFULL
AN
        Amine substituted compounds
TI
        Klein, J. Peter, Vashon, WA, United States
IN
        Underiner, Gail E., Brier, WA, United States
Kumar, Anil M., Seattle, WA, United States
Ridgers, Lance H., Bothell, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 6103730 20000815 <--
PA
PΙ
        US 6103730
                                      19950607 (8)
        US 1995-486264
ΑI
        Continuation of Ser. No. US 1994-217051, filed on 24 Mar 1994, now
RLI
        abandoned
DT
        Utility
        Granted
FS
LN.CNT 1702
        INCLM: 514/263.000
INCL
        INCLS: 514/265.000; 544/268.000; 544/269.000; 544/270.000; 544/271.000;
                 544/272.000
                 514/263.200
NCL
        NCLM:
                 514/151.000; 514/210.210; 514/263.210; 514/263.220; 514/263.230; 514/263.240; 514/263.350; 544/268.000; 544/269.000; 544/270.000; 544/271.000; 544/272.000
        NCLS:
         [7]
IC
        ICM: A61K031-522
         ICS: C07D473-10
         544/268; 544/269; 544/220; 544/271; 544/272; 514/263; 514/265
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 24 OF 78 USPATFULL ON STN
17
        2000:102304 USPATFULL
ΑN
        Therapeutic compounds containing xanthinyl
ΤI
        Klein, J. Peter, Vashon, WA, United States
Leigh, Alistair J., Brier, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
        Kumar, Anil M., Seattle, WA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation) US 6100271 20000808 <--
PA
PΙ
         US 1995-483871
ΑI
                                      19950607 (8)
         Continuation-in-part of Ser. No. US 1994-199368, filed on 18 Feb 1994.
RLI
         now abandoned
         Utility
DT
FS
         Granted
LN.CNT 1986
         INCLM: 514/263.000
INCL
         INCLS: 514/265.000; 544/268.000; 544/269.000; 544/271.000
                  514/263.200
NCL
         NCLM:
                 514/210.210; 514/234.200; 514/263.220; 514/263.230; 514/263.240; 514/263.350; 544/268.000; 544/269.000; 544/271.000
         NCLS:
IC
         ICM: A61K031-522
         ICS: C07D473-10
         544/271; 544/268; 544/269; 514/263; 514/265
FXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 25 OF 78 USPATFULL ON STN
         2000:101856 USPATFULL
ΑN
         Compositions and methods for the treatment and diagnosis of
TI
         cardiovascular disease
         Falb, Dean A., Wellesley, MA, United States
IN
         Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
         corporation)
PΙ
         us 6099823
                                       20000808
         us 1998-126640
                                       19980730 (9)
ΑI
         Continuation-in-part of Ser. No. US 1997-870434, filed on 6 Jun 1997
RLI
         which is a continuation-in-part of Ser. No. US 1997-799910, filed on 13
         Feb 1997
                                  19960216 (60)
PRAI
         US 1996-11787P
         Utility
DT
         Granted
FS
LN.CNT
         5987
         INCLM: 424/009.100
INCL
         INCLS: 536/023.100: 424/009.200; 435/006.000; 435/325.000
         NCLM: 424/009.100
NCL
```

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NCLS: 424/009.200; 435/006.000; 435/325.000; 536/023.100
IC
        [7]
        ICM: C12Q001-68
        ICS: C12N015-85; C12N015-86; C07H021-02; C07H021-04; A61K049-00 435/70.1; 435/325; 435/69.1; 435/6; 435/91.1; 435/91.3; 435/320.1; 435/4; 536/23.1; 536/24.5; 424/9.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 26 OF 78 USPATFULL ON STN
AN
        2000:95042 USPATFULL
        Therapeutic methods employing disulfide derivatives of dithiocarbamates
TT
        and compositions useful therefor
        Lai, Ching-San, Encinitas, CA, United States
ΙN
        Vassilev, Vassil, San Diego, CA, United States
Medinox Inc., San Diego, CA, United States (U.S. corporation)
US 6093743 20000725
PA
        US 6093743
US 1998-103639
PΙ
                                     19980623 (9)
AI
        Utility
DT
FS
        Granted
LN.CNT 2691
        INCLM: 514/599.000
INCL
        INCLS: 514/706.000; 514/707.000; 514/851.000; 514/861.000; 514/863.000;
                 514/866.000; 514/909.000; 514/912.000
                 514/599.000
NCL
        NCLM:
                514/706.000; 514/707.000; 514/851.000; 514/861.000; 514/863.000; 514/866.000; 514/909.000; 514/912.000
        NCLS:
        [7]
IC
        ICM: A61K031-16
        ICS: A61K031-095; A61K031-105
514/599; 514/706; 514/707; 514/851; 514/861; 514/863; 514/866; 514/909;
EXF
        514/912
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 27 OF 78 USPATFULL ON STN
L7
AN
        2000:91735 USPATFULL
        Interferon responsive transcript (IRT-1)
TI
        Autieri, Michael V., Blue Bell, PA, United States
Temple University of the Commonwealth System of Higher Education,
ΙN
PA
        Philadelphia, PA, United States (U.S. corporation)
                                     20000718
PΙ
        us 6090580
ΑI
        us 1998-4171
                                     19980102 (9)
        Utility
DT
        Granted
FS
LN.CNT
        1142
INCL
        INCLM: 435/069.100
        INCLS: 435/252.330; 435/325.000; 435/320.100; 530/350.000; 536/023.100;
                 536/023.500
                 435/069.100
NCL
        NCLM:
                435/252.330; 435/320.100; 435/325.000; 530/350.000; 536/023.100;
        NCLS:
                 536/023.500
IC
        [7]
        ICM: C12P021-06
        ICS: C07H021-04
EXF 530/350; 536/23.5; 435/69.1; 435/252.3; 435/252.33; 435/320.1; 435/325 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 28 OF 78 USPATFULL ON STN
AN
        2000:88304 USPATFULL
TI
        Compositions and methods for the treatment and diagnosis of
        cardiovascular disease
        Falb, Dean A., Wellesley, MA, United States
IN
        Gimbrone, Jr., Michael A., Jamaica Plain, MA, United States
PA
        Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
        corporation)
        Brigham and Women's Hospital, Boston, MA, United States (U.S.
        corporation)
PΙ
        us 6087477
                                     20000711
                                                                                 <--
ΑI
        us 1997-944495
                                     19971006 (8)
        Division of Ser. No. US 1997-799910, filed on 13 Feb 1997
RLI
        US 1996-11787P
                                19960216 (60)
PRAI
        Utility
DT
FS
        Granted
LN.CNT 5589
        INCLM: 530/350.000
INCL
        INCLS: 435/069.100; 435/325.000; 536/023.100
        NCLM: 530/350.000
NCL
```

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NCLS: 435/069.100; 435/325.000; 536/023.100
IC
       [7]
       ICM: C07K016-00
       ICS: C12N015-00
435/325; 435/69.1; 435/6; 536/23.1; 424/185.1; 530/388.24; 530/389.2;
530/350
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 29 OF 78 USPATFULL ON STN
L7
       2000:53742 USPATFULL
AN
       Method of treatment of arterial and venous thromboembolic disorders
TI
       Mousa, Shaker Ahmed, Lincoln University, PA, United States
ΙN
       Dupont Pharmaceuticals, Wilmington, DE, United States (U.S. corporation)
PA
       us 6056958
                                  20000502
ΡI
ΑI
       us 1997-901344
                                  19970728 (8)
       Continuation of Ser. No. US 1994-353419, filed on 9 Dec 1994, now
RLI
       abandoned
DT
       Utility
FS
       Granted
LN.CNT 2186
       INCLM: 424/145.100
INCL
       INCLS: 424/141.100; 424/130.100; 514/002.000
NCL
               424/145.100
               424/130.100; 424/141.100; 514/002.000
       NCLS:
        [7]
IC
       ICM: A61K039-395
EXF 360/32; 360/64; 360/33.1; 360/72.2; 514/2; 386/27; 386/33; 386/37; 386/40-41; 386/109; 386/123; 424/145.1; 424/141.1; 424/130.1 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 30 OF 78 USPATFULL ON STN
L7
       2000:50808 USPATFULL
AΝ
        Compositions and methods for the treatment and diagnosis of
TI
        cardiovascular disease using rchd534 as a target
IN
        Falb, Dean A., Wellesley, MA, United States
        Gimbrone, Jr., Michael A., Jamaica Plain, MA, United States
       Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
        corporation)
        Brigham and Womens's Hospital, Boston, MA, United States (U.S.
        corporation)
                                                                            <--
PΙ
       us 6054558
                                   20000425
       us 1997-925743
                                  19970909 (8)
ΑI
       Division of Ser. No. US 1995-485573, filed on 7 Jun 1995 which is a
RLI
        continuation-in-part of Ser. No. US 1995-386844, filed on 10 Feb 1995
DT
       Utility
        Granted
FS
LN.CNT 5141
        INCLM: 530/350.000
INCL
        INCLS: 536/023.100; 536/024.100; 536/024.300; 435/069.100; 435/320.100;
                435/325.000
               530/350.000
NCL
        NCLM:
               435/069.100; 435/320.100; 435/325.000; 536/023.100; 536/024.100;
        NCLS:
                536/024.300
IC
        [7]
        ICM: C07K016-00
        ICS: C12N015-00
        536/23.1; 536/24.1; 536/24.3; 435/6; 435/810; 435/69.1; 435/7.1; 435/172.3; 435/320.1; 435/325; 435/350; 435/34; 435/52; 435/77; 436/301;
EXF
        436/63; 514/2; 530/388.24; 530/350
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 31 OF 78 USPATFULL on STN
L7
        2000:43949 USPATFULL
AN
ΤI
        Compositions and methods for the treatment and diagnosis of
        cardiovascular disease
        Falb, Dean A., Wellesley, MA, United States
ΙN
        Millennium Pharmaceuticals Inc., Cambridge, MA, United States (U.S.
PA
        corporation)
        us 6048709
                                   20000411
PΙ
        us 1997-826246
                                  19970328 (8)
ΑI
        Division of Ser. No. US 1997-799910, filed on 13 Feb 1997
RLI
        US 1996-11787P
                              19960216 (60)
PRAI
        Utility
DT
        Granted
FS
LN.CNT 5855
        INCLM: 435/069.100
INCL
```

```
INCLS: 435/172.300; 435/252.300; 435/325.000; 435/320.100; 536/023.500;
                536/024.310
NCL
        NCLM:
                435/069.100
                435/252.300; 435/320.100; 435/325.000; 536/023.500; 536/024.310
        NCLS:
IC
        ICM: C12N015-85
        ICS: C12N015-63; C12N015-70; C12N015-12 435/6; 435/69.1; 435/71.2; 435/91.1; 435/172.1; 435/172.3; 435/325;
EXF
        435/375; 435/320.1; 530/350; 536/23.5; 536/25.32
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 32 OF 78 USPATFULL on STN
        2000:43938 USPATFULL
ΑN
        Parallel SELEX.TM.
TI
        Eaton, Bruce, Boulder, CO, United States
Tarasow, Theodore M., Boulder, CO, United States
NeXstar Pharmaceuticals, Inc., Boulder, CO, United States (U.S.
IN
PA
        corporation)
        us 6048698
                                    20000411
PΙ
                                    19980921 (9)
        us 1998-157601
ΑI
        Continuation-in-part of Ser. No. US 1996-618700, filed on 20 Mar 1996,
RLI
        now patented, Pat. No. US 5858660 which is a continuation-in-part of
        Ser. No. US 1994-309245, filed on 20 Sep 1994, now patented, Pat. No. US
        5723289
        Utility
DT
        Granted
FS
LN.CNT 3339
INCL
        INCLM: 435/006.000
        INCLS: 536/025.400
        NCLM:
                435/006.000
NCL
        NCLS:
                536/025.400
IC
        [7]
        ICM: C12Q001-68
        ICS: C07H021-02; C07H021-04
        435/6; 536/25.4
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 33 OF 78 USPATFULL ON STN
L7
        2000:37806 USPATFULL
AN
        Methods for using therapeutic compounds containing xanthinyl
TI
        Klein, J. Peter, Vashon, WA, United States
Leigh, Alistair J., Brier, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
        Kumar, Anil M., Seattle, WA, United States
        Rice, Glenn C., Seattle, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
                                    20000328
PΙ
        us 6043250
        us 1995-472296
                                    19950607 (8)
ΑI
        Continuation-in-part of Ser. No. US 1994-199368, filed on 18 Feb 1994,
RLI
        now abandoned
DT
        Utility
FS
        Granted
LN.CNT
        2052
        INCLM: 514/263.000
INCL
                514/234.200
NCL
        NCLM:
                514/210.210; 514/263.200; 514/263.220; 514/263.230; 514/263.350
        NCLS:
IC
         [7]
        ICM: A61K003-52
EXF
        514/263
L7
      ANSWER 34 OF 78 USPATFULL ON STN
        2000:34403 USPATFULL
AN
        Vascular endothelial growth factor 2
TI
        Hu, Jing-Shan, Sunnyvale, CA, United States
Rosen, Craig A., Laytonsville, MD, United States
IN
        Cao, Liang, South Horizons, Hong Kong
        Human Genome Sciences, Inc., Rockville, MD, United States (U.S.
PA
        corporation)
        us 6040157
                                    20000321
ΡI
        us 1998-42105
                                    19980313 (9)
ΑI
        Continuation-in-part of Ser. No. US 1997-999811, filed on 24 Dec 1997,
RLI
        now patented, Pat. No. US 5932540 which is a continuation-in-part of
        Ser. No. US 1997-824996, filed on 27 Mar 1997 And a continuation-in-part
        of Ser. No. US 1995-465968, filed on 6 Jun 1995 which is a
         continuation-in-part of Ser. No. US 1994-207550, filed on 8 Mar 1994
        Utility
DT
```

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Granted
FS
LN.CNT 5292
       INCLM: 435/069.400
INCL
       INCLS: 435/007.100; 435/325.000; 435/243.000; 435/320.100; 536/023.510;
               530/399.000
               435/069.400
       NCLM:
NCL
               435/007.100; 435/243.000; 435/320.100; 435/325.000; 530/399.000;
       NCLS:
               536/023.510
IC
       Γ71
       ICM: C12N015-18
       ICS: C12N015-63; C12N001-21; C12N005-00
       435/69.4; 435/320.1; 435/325; 435/243; 536/23.51; 530/399
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 35 OF 78 USPATFULL on STN
       2000:12926 USPATFULL
ΑN
       Compositions and methods for the treatment and diagnosis of
TI
       cardiovascular disease using rchd523 as a target
       Falb, Dean A., Wellesley, MA, United States
IN
       Gimbrone, Jr., Michael A., Jamaica Plain, MA, United States
PA
       Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
       corporation)
       Brigham and Women's Hospital, Boston, MA, United States (U.S.
       corporation)
       us 6020463
                                                                         <--
PΙ
                                 20000201
       us 1997-944423
ΑI
                                 19971006 (8)
       Division of Ser. No. US 1996-599654, filed on 9 Feb 1996, now patented, Pat. No. US 5882925 which is a continuation-in-part of Ser. No. US
RLI
       1995-485573, filed on 7 Jun 1995, now patented, Pat. No. US 5968770
       which is a continuation-in-part of Ser. No. US 1995-386844, filed on 10
       Feb 1995
       Utility
DT
FS
       Granted
       5972
LN.CNT
INCL
       INCLM: 530/350.000
        INCLS: 435/069.100; 435/320.100; 435/325.000; 536/023.100
NCL
               530/350.000
               435/069.100; 435/320.100; 435/325.000; 536/023.100
       NCLS:
        [6]
IC
        ICM: C07K016-00
        ICS: C12N015-00
        435/320.1; 435/325; 435/69.1; 435/6; 536/23.1; 530/350
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 36 OF 78 USPATFULL on STN
L7
        2000:12800 USPATFULL
AN
TI
        Electronegative-substituted long chain xanthine compounds
       Leigh, Alistair J., Brier, WA, United States
IN
       Michnick, John, Seattle, WA, United States
       Kumar, Anil M., Seattle, WA, United States
Klein, J. Peter, Vashon, WA, United States
        Underiner, Gail, Malvern, PA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 6020337 20000201 <--
PA
PΙ
       US 1997-950810
                                 19970916 (8)
ΑI
       Continuation-in-part of Ser. No. US 1993-42946, filed on 5 Apr 1993, now
RLI
        patented, Pat. No. US 5670506 And a continuation-in-part of Ser. No. US
        1997-910579, filed on 26 Jul 1997
DT
        Utility
FS
        Granted
LN.CNT 1376
INCL
        INCLM: 514/258.000
               514/263.000; 544/267.000; 544/272.000; 544/277.000
        INCLS:
NCL
               514/263.340
        NCLM:
               514/210.210; 514/263.360; 544/267.000; 544/272.000; 544/277.000
        NCLS:
IC
        [6]
        ICM: A61K031-52
        ICS: C07D473-00
        514/258; 544/267; 544/272; 544/277
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 37 OF 78 USPATFULL ON STN
L7
        2000:10014 USPATFULL
ΑN
        Compositions and methods for the treatment and diagnosis of
TI
        cardiovascular disease using rchd528 as a target
        Falb, Dean A., Wellesley, MA, United States
IN
```

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Gimbrone, Jr., Michael A., Jamaica Plain, MA, United States
       Millenium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
       corporation)
       Brigham and Women's Hospital, Boston, MA, United States (U.S.
       corporation)
       US 6018025
                                                                          <--
                                  20000125
PΙ
       US 1997-944868
                                  19971006 (8)
ΑI
       Division of Ser. No. US 1996-599654, filed on 9 Feb 1996, now patented,
RLI
       Pat. No. US 5882925 which is a continuation-in-part of Ser. No. US
       1995-485573, filed on 7 Jun 1995 which is a continuation-in-part of Ser.
       No. US 1995-386844, filed on 10 Feb 1995
DT
       Utility
       Granted
FS
LN.CNT 6133
INCL
       INCLM: 530/350.000
                             530/326.000; 536/023.100; 536/023.500; 435/069.100;
               530/324.000;
               435/320.100;
                            435/325.000
               530/350.000
NCL
       NCLM:
               435/069.100; 435/320.100; 435/325.000; 530/324.000; 530/326.000;
       NCLS:
               536/023.100; 536/023.500
       [6]
IC
       ICM: C07K016-00
       ICS: C12N015-00
       536/23.1; 536/24.1; 536/24.3; 536/23.5; 435/6; 435/810; 435/69.1; 435/7.1; 435/172.3; 435/320.1; 435/325; 436/501; 436/63; 935/32; 935/52;
EXF
       935/77; 530/350; 530/325; 530/326
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 38 OF 78 USPATFULL on STN
L7
       2000:4941 USPATFULL
AN
          ***VEGF*** .sub.145 expression vectors
ΤI
       Neufeld, Gera, Haifa, Israel
ΙN
       Keshet, Eli, Kiryat Yam, Israel
       Vlodavsky, Ísrael, Mevaseret Zion, Israel
Poltorak, Zoya, Jerusalem, Israel
       Technion Research & Development Co. Ltd., Haifa, Israel (non-U.S.
       corporation)
                                  20000111
                                                                          <--
PΙ
       us 6013780
       US 1997-784551
                                  19970121 (8)
AΙ
       US 1996-25537P
                             19960906 (60)
PRAI
       Utility
DT
FS
       Granted
LN.CNT 2158
       INCLM: 536/023.100
INCL
       INCLS: 435/320.100
               536/023.100
NCL
       NCLM:
       NCLS:
               435/320.100
IC
        [6]
       ICM: C07H021-04
       ICS: C12N015-11; C12N015-63
514/44; 435/172.3; 435/320.1; 536/23.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 39 OF 78 USPATFULL ON STN
        1999:151195 USPATFULL
AN
       GATA-6 transcription factor: compositions and methods
TI
       Walsh, Kenneth, Carlisle, MA, United States
IN
PA
       St. Elizabeth's Medical Center, Boston, MA, United States (U.S.
        corporation)
                                  19991123
PΙ
       us 5990092
                                                                          <--
       US 1997-927394
                                  19970827 (8)
ΑI
DT
       Utility
FS
        Granted
LN.CNT 2449
INCL
        INCLM: 514/044.000
        INCLS: 435/320.100; 536/023.500
               514/044.000
NCL
       NCLM:
       NCLS:
               435/320.100; 536/023.500
IC
        [6]
        ICM: A61K048-00
        ICS: C12N015-12; C12N015-85
        435/320.1; 435/375; 435/377; 514/44; 536/23.5
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 40 OF 78 USPATFULL ON STN
ı 7
```

1999:128386 USPATFULL

AN

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Compositions and methods for the treatment and diagnosis of
ΤI
        cardiovascular disease using rchd523 as a target
       Falb, Dean A., Wellesley, MA, United States
Gimbrone, Jr., Michael A., Jamaica Plain, MA, United States
Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
IN
PA
        corporation)
PΙ
        US 5968770
                                    19991019
        US 1995-485573
                                    19950607 (8)
ΑI
        Continuation-in-part of Ser. No. US 1995-386844, filed on 10 Feb 1995
RLI
DT
        Granted
FS
LN.CNT 5019
        INCLM: 435/069.100
INCL
        INCLS: 435/006.000; 435/007.100; 435/320.100; 435/325.000; 435/455.000; 514/044.000; 536/023.100; 536/024.100; 536/024.300
NCL
        NCLM:
                435/069.100
                435/006.000; 435/007.100; 435/320.100; 435/325.000; 435/455.000;
        NCLS:
                514/044.000; 536/023.100; 536/024.100; 536/024.300
IC
        [6]
        ICM: C12N001-21
        536/23.1; 536/24.1; 536/24.3; 435/6; 435/69.1; 435/7.1; 435/320.1;
EXF
        435/325; 435/455; 514/44; 436/501; 436/63
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 41 OF 78 USPATFULL ON STN 1999:96407 USPATFULL
L7
ΑN
        Pulsed administration of compositions for the treatment of blood
TI
        disorders
        Perrine, Susan P., 27 Harding Ave., Braintree, MA, United States
ΙN
        us 5939456
                                    19990817
PΙ
                                    19960726 (8)
ΑI
        us 1996-687670
        Utility
DT
FS
        Granted
LN.CNT 2147
INCL
        INCLM: 514/554.000
        INCLS: 514/538.000; 514/546.000; 514/563.000; 514/568.000; 514/576.000; 514/578.000; 514/629.000
NCL
        NCLM:
                514/554.000
                514/538.000; 514/546.000; 514/563.000; 514/568.000; 514/576.000; 514/578.000; 514/629.000
        NCLS:
IC
        [6]
        ICM: A61K031-205
        ICS: A61K031-19; A61K031-22
EXF
        514/576; 514/578; 514/563; 514/568; 514/538; 514/629; 514/546; 514/554
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 42 OF 78 USPATFULL ON STN
        1999:92656 USPATFULL
ΑN
        Compositions and methods for modulating growth of a tissue in a mammal
TI
        Weisz, Paul B., State College, PA, United States
Trustees of the University of Pennsylvania, Philadelphia, PA, United
IN
PA
        States (U.S. corporation)
        us 5935940
                                    19990810
PΙ
        us 1997-906500
                                    19970805 (8)
ΑI
        Division of Ser. No. US 1994-345011, filed on 23 Nov 1994, now patented,
RLI
        Pat. No. US 5658894 which is a continuation of Ser. No. US 1992-900592,
        filed on 18 Jun 1992, now abandoned And a continuation-in-part of Ser.
        No. US 1991-790320, filed on 12 Nov 1991, now abandoned which is a
        continuation of Ser. No. US 1991-691168, filed on 24 Apr 1991, now
        abandoned which is a continuation of Ser. No. US 1989-397559, filed on
        23 Aug 1989, now abandoned , said Ser. No. US 900592 which is a continuation-in-part of Ser. No. US 1990-480407, filed on 15 Feb 1990,
        now patented, Pat. No. US 5183809
DT
        Utility
FS
        Granted
LN.CNT 1497
INCL
        INCLM: 514/058.000
        INCLS: 514/021.000; 530/810.000; 530/812.000; 530/813.000
                514/058.000
NCL
        NCLM:
        NCLS:
                514/021.000; 530/810.000; 530/812.000; 530/813.000
IC
        [6]
        ICM: A61K031-715
        ICS: A61K038-00
        514/58; 514/21; 530/810; 530/812; 530/813
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
```

```
ANSWER 43 OF 78 USPATFULL ON STN
L7
        1999:89116 USPATFULL
AN
ΤI
        Vascular endothelial growth factor 2
        Hu, Jing-Shan, Sunnyvale, CA, United States
IN
        Rosen, Craig A., Laytonsville, MD, United States
        Cao, Liang, Hong Kong, Hong Kong
PA
        Human Genome Sciences, Inc., Rockville, MD, United States (U.S.
        corporation)
                                     19990803
PΙ
        US 5932540
        us 1997-999811
                                     19971224 (8)
ΑI
        Continuation-in-part of Ser. No. US 1994-207550, filed on 8 Mar 1994,
RLI
        now abandoned And Ser. No. US 1995-465968, filed on 6 Jun 1995
        Utility
DT
FS
        Granted
LN.CNT 2605
        INCLM: 514/002.000
INCL
        INCLS: 530/326.000; 530/399.000; 530/402.000
                 514/002.000
NCL
        NCLM:
                 530/326.000; 530/399.000; 530/402.000
        NCLS:
        [6]
TC
        ICM: A61K038-14
        ICS: C07K014-475
        514/2; 514/12; 530/399; 530/326; 530/402
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 44 OF 78 USPATFULL on STN
AN
        1999:56471 USPATFULL
        Methods of modulating tissue growth and regeneration
TI
        Herrmann, Howard C., Bryn Mawr, PA, United States
IN
        Barnathan, Elliot, Havertown, PA, United States
        Weisz, Paul B., State College, PA, United States
The Trustees of the University of Pennsylvania, Philadelphia, PA, United
PA
        States (U.S. corporation)
        US 5902799
                                     19990511
PΙ
        us 1997-906501
                                     19970805 (8)
ΑI
        Division of Ser. No. US 1994-345011, filed on 23 Nov 1994, now patented, Pat. No. US 5658894 which is a continuation of Ser. No. US 1992-900592,
RLI
        filed on 18 Jun 1992, now abandoned And a continuation-in-part of Ser. No. US 1991-790320, filed on 12 Nov 1991, now abandoned which is a continuation of Ser. No. US 1991-691168, filed on 24 Apr 1991, now
        abandoned which is a continuation of Ser. No. US 1989-397559, filed on
        23 Aug 1989, now abandoned , said Ser. No. US 900592 which is a continuation-in-part of Ser. No. US 1990-480407, filed on 15 Feb 1990,
        now_patented, Pat. No. US 5183809
DT
        Utility
FS
        Granted
LN.CNT 1703
INCL
        INCLM: 514/058.000
        INCLS: 514/021.000; 530/810.000; 530/813.000; 530/817.000
NCL
        NCLM:
                 514/058.000
                 514/021.000; 530/810.000; 530/813.000; 530/817.000
        NCLS:
IC
        [6]
        ICM: A61K031-715
        ICS: A61K031-735
        514/58; 514/21; 514/56; 530/810; 530/812; 530/813; 530/817
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 45 OF 78 USPATFULL ON STN
AN
        1999:40428 USPATFULL
TI
        Substituted amino alkyl compounds
IN
        Klein, J. Peter, Vashon Island, WA, United States
        Underiner, Gail E., Brier, WA, United States
Leigh, Alistair J., Brier, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 5889011 19990330 <--
PA
PΙ
                                     19970627 (8)
        us 1997-884037
ΑI
        Continuation of Ser. No. US 1993-149681, filed on 9 Nov 1993, now
RLI
        abandoned which is a continuation-in-part of Ser. No. US 1992-973804,
        filed on 9 Nov 1992, now patented, Pat. No. US 5340813
DT
        Utility
FS
        Granted
LN.CNT 1351
        INCLM: 514/263.000
INCL
        INCLS: 514/261.000; 544/267.000; 544/264.000; 544/265.000
NCL
        NCLM:
                 514/263.350
                 544/264.000; 544/265.000; 544/267.000
        NCLS:
```

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[6]
IC
       ICM: C07D473-00
       ICS: A61K031-52
544/257; 544/263; 544/285; 544/287; 514/263; 514/261
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 46 OF 78 USPATFULL ON STN
       1999:33831 USPATFULL
ΑN
TI
       Compositions and method for the treatment and diagnosis of
       cardiovascular disease using rchd502 as a target
       Falb, Dean A., Wellesley, MA, United States
ΙN
       Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
       corporation)
US 5882925
US 1996-599654
PΙ
                                 19990316
                                                                         <--
                                 19960209 (8)
ΑI
       Continuation-in-part of Ser. No. US 1995-485573, filed on 7 Jun 1995
RLI
       which is a continuation-in-part of Ser. No. US 1995-386844, filed on 10
       Feb 1995
       Utility
DT
       Granted
FS
LN.CNT 5758
INCL
       INCLM: 435/325.000
       INCLS: 536/023.100; 536/024.100; 536/024.300; 435/006.000; 435/069.100;
               435/320.100; 435/455.000
       NCLM:
               435/325.000
NCL
               435/006.000; 435/069.100; 435/320.100; 435/455.000; 536/023.100;
       NCLS:
               536/024.100; 536/024.300
IC
       [6]
       ĪCM: C12N015-12
       536/23.1; 536/24.1; 536/24.3; 435/6; 435/69.1; 435/325; 435/320.1;
EXF
       435/455
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 47 OF 78 USPATFULL on STN
ΑN
       1999:24638 USPATFULL
       Compositions and methods for modulating growth of a tissue in a mammal
TI
       Herrmann, Howard C., Bryn Mawr, PA, United States
TN
       Barnathan, Elliot, Havertown, PA, United States
       Weisz, Paúl B., State Collegé, PÁ, United States
The Trustees of the University of Pennsylvania, Philadelphia, PA, United
PA
       States (U.S. corporation)
       us 5874419
                                 19990223
PΙ
       US 1997-905612
                                 19970804 (8)
ΑI
RLI
       Division of Ser. No. US 1994-345011, filed on 23 Nov 1994, now patented,
       Pat. No. US 5658894 which is a continuation of Ser. No. US 1992-900592,
       filed on 18 Jun 1992, now abandoned And a continuation-in-part of Ser.
       No. US 1991-790320, filed on 12 Nov 1991, now abandoned which is a
       continuation-in-part of Ser. No. US 1991-691168, filed on 24 Apr 1991,
       now abandoned which is a continuation of Ser. No. US 1989-397559, filed on 23 Aug 1989, now abandoned , said Ser. No. US 20 -900592 which is a
       continuation-in-part of Ser. No. US 1990-480407, filed on 15 Feb 1990,
       now patented, Pat. No. US 5183809, issued on 2 Feb 1993
DT
       Utility
       Granted
FS
LN.CNT 1482
INCL
       INCLM: 514/058.000
       INCLS: 514/021.000; 514/023.000; 514/054.000; 514/060.000; 514/769.000;
               424/652.000; 424/682.000; 424/617.000; 536/103.000
NCL
               514/058.000
       NCLM:
               424/617.000; 424/652.000; 424/682.000; 514/021.000; 514/023.000;
       NCLS:
               514/054.000; 514/060.000; 514/769.000; 536/103.000
IC
       [6]
       ICM: A61K031-735
       ICS: A61K047-02; C08B037-16
                        514/54; 514/58; 514/60; 514/769; 536/103; 424/652;
       514/21; 514/23;
EXF
       424/682; 424/617
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 48 OF 78 USPATFULL ON STN
L7
       1999:18711 USPATFULL
ΑN
TI
       Adenoviral-mediated gene transfer to adipocytes
       Crystal, Ronald G., Potomac, MD, United States
IN
       Magovern, Christopher J., New York, NY, United States
PA
       Cornell Research Foundation, Inc., Ithaca, NY, United States (U.S.
       corporation)
       US 5869037
                                 19990209
PΙ
                                                                         <--
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19960626 (8)
       us 1996-672461
ΑI
       Utility
DT
FS
       Granted
LN.CNT 1452
       INCLM: 424/093.200
INCL
       INCLS: 424/093.700; 424/093.210; 435/325.000; 435/320.100; 435/172.300;
               514/044.000
               424/093.200
NCL
       NCLM:
               424/093.210; 424/093.700; 435/320.100; 435/325.000; 435/456.000;
       NCLS:
               514/044.000
        [6]
TC
       ICM: A61K035-12
       ICS: A61K048-00; C12N015-09; C12N015-86
       514/2; 514/44; 435/172.3; 435/320.1; 435/252.3; 435/325.1; 424/93.21;
EXF
        424/93.7; 424/93.2; 536/24.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 49 OF 78 USPATFULL on STN
                   USPATFULL
        1999:4647
ΑN
       Fas ligand compositions for treatment of proliferative disorders
TI
       walsh, Kenneth, Carlisle, MA, United States
TN
       St. Elizabeth's Medical Center, Boston, MA, United States (U.S.
PA
        corporation)
                                  19990112
       us 5858990
                                                                           <--
PΙ
                                  19970304 (8)
       us 1997-810453
ΑI
       Utility
DT
FS
        Granted
LN.CNT 3038
INCL
        INCLM: 514/044.000
        INCLS: 435/006.000; 435/172.100; 435/320.100; 435/069.100; 435/375.000;
               435/377.000
               514/044.000
NCL
        NCLM:
               435/006.000; 435/069.100; 435/320.100; 435/375.000; 435/377.000
        NCLS:
        [6]
IC
        ICM: A61K048-00
        ICS: C12N015-11
        435/6; 435/172.1; 435/172.3; 435/320.1; 435/325; 435/69.1; 435/31.1; 435/375; 435/377; 536/23.1; 536/23.5; 514/2; 514/44
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 50 OF 78 USPATFULL ON STN
L7
        1999:4328 USPATFULL
AN
        Parallel selex
TI
        Eaton, Bruce, Boulder, CO, United States
ΙN
        Gold, Larry, Boulder, CO, United States
        Nexstar Pharmaceuticlas, Inc., Boulder, CO, United States (U.S.
PA
        corporation)
PΙ
        us 5858660
                                  19990112
                                  19960320 (8)
        us 1996-618700
ΑI
        Continuation-in-part of Ser. No. US 1994-309245, filed on 20 Sep 1994,
RLI
        now patented, Pat. No. US 5723289
DT
        Utility
FS
        Granted
LN.CNT
        3236
INCL
        INCLM: 435/006.000
        INCLS: 435/091.200; 536/025.400; 536/022.100; 935/077.000; 935/078.000
NCL
                435/006.000
        NCLM:
        NCLS:
               435/091.200; 536/022.100; 536/025.400
IC
        [6]
        ICM: C12Q001-68
        ICS: C120019-34; C07H021-02; C07H021-04
435/6; 435/91.2; 536/22.1; 536/25.4; 935/77; 935/78
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 51 OF 78 USPATFULL ON STN
        1998:157185 USPATFULL
ΑN
        Compositions and methods for the treatment and diagnosis of
TI
        cardiovascular using RCHD528 as a target
        Falb, Dean A., Massachusetts, MA, United States
IN
        Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
        corporation)
        us 5849578
                                  19981215
PΙ
                                                                           <--
                                  19960315 (8)
        us 1996-616844
ΑI
        Division of Ser. No. US 1996-599654, filed on 9 Feb 1996 which is a continuation-in-part of Ser. No. US 1995-458873, filed on 7 Jun 1995
RLI
        which is a continuation-in-part of Ser. No. US 1995-386844, filed on 10
```

```
Feb 1995
DT
         Utility
FS
         Granted
         5753
LN.CNT
          INCLM: 435/325.000
INCL
                   536/023.100; 536/024.100; 536/024.300; 435/006.000; 435/069.100;
         INCLS:
                    435/320.100; 435/455.000
NCL
         NCLM:
                    435/325.000
                    435/006.000; 435/069.100; 435/320.100; 435/455.000; 536/023.100;
         NCLS:
                    536/024.100: 536/024.300
IC
          [6]
          ICM: C12N015-12
         536/23.1; 536/24.1; 536/24.3; 435/6; 435/69.1; 435/7.1; 435/325; 435/320.1; 435/455; 436/201; 436/63; 514/44
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
       ANSWER 52 OF 78 USPATFULL ON STN
          1998:144102 USPATFULL
ΑN
          Amino-alcohol substituted cyclic compounds
TI
         Kumar, Anil M., Seattle, WA, United States
ΙN
         Michnick, John, Seattle, WA, United States
Underiner, Gail E., Brier, WA, United States
          Klein, J. Peter, Vashon Island, WA, United States
          Rice, Glenn C., Seattle, WA, United States
          Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
PΙ
                                            19981117
          us 5837703
          US 1993-152650
                                            19931112 (8)
ΑI
RLI
          Continuation-in-part of Ser. No. US 1993-40820, filed on 31 Mar 1993,
          now abandoned
DT
          Utility
          Granted
FS
LN.CNT 2596
INCL
          INCLM: 514/183.000
          INCLS: 514/211.000; 514/228.800; 514/241.000; 514/242.000; 514/249.000;
                    514/256.000; 514/259.000; 514/263.000; 514/270.000; 514/274.000;
                    514/309.000; 514/312.000; 514/315.000; 514/348.000; 514/357.000;
                    514/374.000; 514/400.000; 514/425.000; 514/427.000; 540/467.000; 540/544.000; 544/216.000; 544/257.000; 544/272.000; 544/286.000; 544/301.000; 544/311.000; 544/335.000; 546/096.000; 546/141.000; 546/142.000; 546/157.000; 546/246.000; 546/296.000; 546/334.000; 548/215.000; 548/340.100; 548/485.000; 548/546.000; 548/561.000
NCL
          NCLM:
                    514/183.000
                                      514/228.800; 514/241.000; 514/242.000; 514/249.000; 514/266.200; 514/266.300; 514/270.000; 514/274.000;
                    514/211.150;
          NCLS:
                    514/256.000;
                                      514/266.200;
                    514/309.000; 514/312.000; 514/315.000; 514/348.000; 514/357.000;
                    514/374.000; 514/400.000; 514/425.000; 514/427.000; 540/467.000;
                    540/544.000; 544/216.000; 544/257.000; 544/272.000; 544/286.000;
                    544/301.000; 544/311.000; 544/335.000; 546/096.000; 546/141.000; 546/142.000; 546/157.000; 546/246.000; 546/296.000; 546/334.000; 548/215.000; 548/340.100; 548/485.000; 548/546.000; 548/561.000
IC
          [6]
          ICM: A61K031-55
         ICS: A61K031-53

ICS: A61K031-515; A61K031-445; A61K031-52

544/276; 544/272; 544/216; 544/257; 544/285; 544/286; 544/301; 544/311;

544/335; 514/263; 514/183; 514/211; 514/228.8; 514/241; 514/242;

514/249; 514/256; 514/259; 514/270; 514/274; 514/309; 514/312; 514/315;

514/348; 514/357; 514/374; 514/400; 514/418; 514/425; 514/427; 540/467;

540/544; 546/96; 546/141; 546/142; 546/157; 546/246; 546/296; 546/334;
EXF
          548/215; 548/340.1; 548/485; 548/546; 548/561
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
       ANSWER 53 OF 78 USPATFULL on STN
ΑN
          1998:138691 USPATFULL
TI
          Compositions and methods using rchd534, a gene uregulated by shear
ΙN
          Falb, Dean, Wellesley, MA, United States
          Millennium Pharmaceuticals Inc., Cambridge, MA. United States (U.S.
PA
          corporation)
PΙ
          us 5834248
                                            19981110
                                            19950607 (8)
          us 1995-480994
ΑI
RLI
          Division of Ser. No. US 1995-485573, filed on 7 Jun 1995 And a
          continuation-in-part of Ser. No. US 1995-386844, filed on 10 Feb 1995
DT
          Utility
          Granted
FS
LN.CNT 4877
```

INCLM: 435/070.100

INCL

```
INCLS: 435/325.000; 435/172.300; 435/320.100; 536/023.100; 536/023.500
                435/070.100
NCL
        NCLM:
                435/320.100; 435/325.000; 536/023.100; 536/023.500
        NCLS:
        [6]
IC
        ICM: C12N015-00
        ICS: C07H021-00
EXF
        514/44; 424/93.1; 536/23.1; 536/23.5; 435/325
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 54 OF 78 USPATFULL ON STN
L7
ΑN
        1998:128265 USPATFULL
ΤI
        Substituted amino alcohol compounds
       Klein, J. Peter, Vashon, WA, United States
Underiner, Gail E., Brier, WA, United States
Kumar, Anil M., Seattle, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 5824677
19981020
---
IN
PA
PΙ
        US 1995-474816
ΑI
                                    19950607 (8)
RLI
        Division of Ser. No. US 1994-303842, filed on 8 Sep 1994, now patented,
        Pat. No. US 5641783 which is a continuation-in-part of Ser. No. US
        1993-152650, filed on 12 Nov 1993, now patented, Pat. No. US 5801181 And
        Ser. No. US 1993-164081, filed on 8 Dec 1993, now patented, Pat. No. US
        5470878 , said Ser. No. US
                                         -152650 And Ser. No. US
                                                                       -164081 , each
                        - which is a continuation-in-part of Ser. No. US
        1993-40820, filed on 31 Mar 1993, now abandoned
DT
        Utility
FS
        Granted
        3136
LN.CNT
        INCLM: 514/222.500
INCL
        INCLS: 514/223.500; 514/224.500; 514/226.800; 514/227.500; 514/228.800;
                514/229.200; 514/230.500;
                                              514/230.800;
                                                             514/237.800;
                                                                            514/248.000;
                514/249.000; 514/255.000; 514/258.000; 514/274.000;
                                                                            514/301.000;
                514/303.000; 514/311.000; 514/351.000; 514/360.000;
                                                                            514/361.000;
                514/362.000; 514/363.000; 514/364.000; 514/365.000;
                                                                            514/367.000;
                514/372.000; 514/373.000; 514/374.000; 514/375.000;
                                                                            514/376.000;
                514/378.000; 514/379.000; 514/380.000; 514/387.000; 514/415.000; 514/418.000; 514/424.000; 514/425.000;
                                                                            514/395.000;
                                                                            514/433.000;
                                             514/438.000;
514/438.000;
549/014.000;
544/002.000;
544/063.000;
544/091.000;
                              514/432.000;
346/300.000;
549/368.000;
                514/452.000;
                                                             346/113.000;
                                                                            346/114.000;
                346/164.000;
                                                             549/050.000;
                                                                            549/075.000;
                                                             544/003.000;
                549/367.000;
                                                                            544/005.000
                               544/053.000;
                544/008.000;
                                                             544/065.000;
                                                                            544/066.000
                                                             544/127.000;
                                                                            544/128.000
                544/067.000;
                               544/090.000;
                544/162.000;
                               544/215.000;
                                              544/219.000;
                                                             544/229.000;
                                                                            544/235.000
                               544/255.000;
                                              544/278.000;
                                                                            544/353.000;
                544/237.000;
                                                             544/311.000;
                                              548/125.000;
                544/385.000;
                              548/123.000;
                                                             548/131.000;
                                                                            548/134.000:
                                              548/153.000;
                                                             548/174.000;
                548/143.000; 548/146.000;
                                                                           548/207.000;
                548/214.000;
                               548/215.000; 548/217.000;
                                                             548/221.000; 548/228.000;
                              548/237.000; 548/240.000;
                548/229.000;
                                                             548/241.000; 548/243.000;
                548/247.000;
                               548/267.200; 548/303.700; 548/307.100; 548/453.000;
                548/486.000; 548/543.000; 548/546.000
NCL
                514/222.500
        NCLM:
                514/223.500;
                               514/224.500; 514/226.800; 514/227.500; 514/228.800;
        NCLS:
                514/229.200;
                               514/230.500;
                                              514/230.800;
                                                             514/237.800;
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                                              514/260.100;
                514/249.000;
                               514/255.020;
                                                             514/274.000;
                                                                            514/301.000;
                               514/311.000;
                                              514/351.000;
                                                             514/360.000;
                514/303.000;
                                                                            514/361.000
                                                            514/365.000;
                514/362.000;
                               514/363.000;
                                              514/364.000;
                                                                            514/367.000
                              514/373.000; 514/374.000; 514/375.000; 514/376.000;
                514/372.000;
                                             514/380.000; 514/387.000; 514/395.000;
                514/378.000;
                               514/379.000;
                               514/418.000;
                514/415.000;
                                             514/424.000; 514/425.000;
                                                                           514/432.000;
                               514/438.000;
                                             514/452.000; 544/002.000;
                514/433.000;
                                                                           544/003.000;
                              544/008.000;
544/067.000;
544/162.000;
544/237.000;
                544/005.000;
                                             544/053.000;
                                                            544/063.000;
                                                                           544/065.000;
                                                            544/091.000;
                544/066.000;
                                              544/090.000;
                                                                            544/127.000
                                              544/215.000;
544/255.000;
                544/128.000;
                                                             544/219.000;
                                                                            544/229.000
                                                             544/278.000;
                544/235.000;
                                                                            544/311.000;
                               544/385.000;
                                              546/113.000;
                                                             546/114.000;
                544/353.000;
                                                                            546/164.000
                546/300.000;
                               548/123.000;
                                              548/125.000;
                                                             548/131.000:
                                                                            548/134.000
                                                             548/174.000;
                548/143.000:
                               548/146.000:
                                              548/153.000;
                                                                            548/207.000
                                                             548/221.000;
                               548/215.000;
                548/214.000;
                                              548/217.000:
                                                                            548/228.000
                548/229.000;
                               548/237.000;
                                             548/240.000;
                                                             548/241.000;
                                                                           548/243.000;
                548/247.000;
                              548/267.200; 548/303.700;
                                                            548/307.100;
                                                                           548/453.000;
                548/486.000; 548/543.000; 548/546.000
549/075.000; 549/367.000; 549/368.000
                               548/543.000; 548/546.000; 549/014.000; 549/050.000;
IC
        [6]
        ICM: A61K031-385
        ICS: A61K031-445; A61K031-47; A61K031-505
        549/75; 549/50; 549/14; 549/367; 549/368; 514/432; 514/438; 514/222.5;
EXF
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514/223.5; 514/224.5; 514/226.8; 514/227.5; 514/228.8; 514/229.2; 514/230.5; 514/230.8; 514/237.8; 514/248; 514/249; 514/255; 514/258; 514/274; 514/301; 514/303; 514/311; 514/351; 514/360; 514/361; 514/362; 514/363; 514/364; 514/365; 514/367; 514/372; 514/373; 514/374; 514/375; 514/376; 514/378; 514/379; 514/380; 514/387; 514/395; 514/415; 514/418; 514/424; 514/425; 514/433; 514/452; 544/2; 544/3; 544/5; 544/8; 544/53; 544/63; 544/65; 544/66; 544/67; 544/90; 544/91; 544/127; 544/128; 544/162; 544/215; 544/219; 544/229; 544/235; 544/237; 544/255; 544/278; 544/311; 544/353; 544/385; 546/113; 546/114; 546/164; 546/300; 548/123; 548/125; 548/131; 548/134; 548/145; 548/146; 548/153; 548/174; 548/207
          548/125; 548/131; 548/134; 548/145; 548/146; 548/153; 548/174; 548/207; 548/214; 548/215; 548/217; 548/221; 548/228; 548/229; 548/237; 548/240;
          548/241; 548/243; 548/247; 548/267.2; 548/303.7; 548/307.1; 548/453;
          548/486; 548/543; 548/546
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
       ANSWER 55 OF 78 USPATFULL ON STN
          1998:122413 USPATFULL
AN
          Substituted amino alkyl compounds
ΤI
          Klein, J. Peter, Vashon Island, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
          Leigh, Alistair J., Brier, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
                                             19981006
PI
          us 5817662
          US 1995-468656
                                             19950606 (8)
ΑI
          Division of Ser. No. US 1993-149681, filed on 9 Nov 1993, now abandoned
RLI
          which is a continuation-in-part of Ser. No. US 1992-973804, filed on 9
          Nov 1992, now patented, Pat. No. US 5340813
DT
          Utility
FS
          Granted
          1358
LN.CNT
          INCLM: 514/263.000
INCL
          INCLS: 424/824.000; 424/825.000; 424/885.000; 424/921.000
NCL
                    514/263.350
          NCLM:
                    424/824.000; 424/825.000
          NCLS:
IC
          [6]
          ICM: A61K031-52
          514/397; 514/263; 424/824; 424/825; 424/885; 424/921
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
       ANSWER 56 OF 78 USPATFULL on STN
          1998:111942 USPATFULL
ΑN
ΤI
          Therapeutic compounds containing pyrimidinyl moieties
          Klein, J. Peter, Vashon, WA, United States
Leigh, Alistair J., Brier, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
          Kumar, Anil M., Seattle, WA, United States
          Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation) US 5807862 19980915 <--
PA
PΙ
ΑI
          us 1995-478112
                                             19950607 (8)
          Continuation-in-part of Ser. No. US 1994-199368, filed on 18 Feb 1994,
RLI
          now abandoned
DT
          Utility
FS
          Granted
LN.CNT
          2190
INCL
          INCLM: 514/269.000
          INCLS: 544/309.000; 544/310.000; 544/311.000; 544/312.000
NCL
                     514/269.000
          NCLM:
          NCLS:
                    544/309.000; 544/310.000; 544/311.000; 544/312.000
          [6]
IC
          ICM: A61K031-505
          ICS: C07D239-54
          514/269; 514/274; 544/309; 544/310; 544/311; 544/312
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
       ANSWER 57 OF 78 USPATFULL on STN
          1998:111941 USPATFULL
ΑN
          Amine substituted xanthinyl compounds
TI
          Klein, J. Peter, Vashon, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
          Kumar, Anil M., Seattle, WA, United States
          Ridgers, Lance H., Bothell, WA, United States
          Rice, Glenn C., Seattle, WA, United States
          Leung, David W., Mercer Island, WA, United States
          Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
          US 5807861
US 1995-476911
                                             19980915
PΙ
                                             19950607 (8)
ΑI
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Continuation-in-part of Ser. No. US 1994-217051, filed on 24 Mar 1994,
RLI
        now abandoned
DT
        Utility
FS
        Granted
LN.CNT
       1713
        INCLM: 514/263.000
INCL
                514/263.350
        NCLM:
NCL
                514/081.000; 514/151.000; 514/210.210; 514/263.200; 514/263.220;
        NCLS:
                514/263.230
IC
        [6]
        ICM: A61K031-52
        514/263
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 58 OF 78 USPATFULL on STN
L7
AN
        1998:108415 USPATFULL
        Therapeutic compounds containing a monocyclic five- to six- membered
TI
        ring structure having one to two nitrogen atoms
        Underiner, Gail E., Brier, WA, United States
IN
        Porubek, David, Seattle, WA, United States
        Klein, J. Peter, Vashon Island, WA, United States
        Woodson, Paul, Edmonds, WA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 5804584 19980908 <--
PA
PΙ
        US 1995-468659 19950606 (8)
Division of Ser. No. US 1993-153256, filed on 16 Nov 1993, now abandoned
AI
RLI
        which is a continuation-in-part of Ser. No. US 1992-976353, filed on 16
        Nov 1992, now patented, Pat. No. US 5473070
DT
        Utility
        Granted
FS
LN.CNT 1554
        INCLM: 514/269.000
INCL
        INCLS: 544/298.000; 544/242.000; 544/301.000; 544/302.000; 514/256.000
                514/269.000
NCL
        NCLM:
        NCLS:
                514/256.000; 544/242.000; 544/298.000; 544/301.000; 544/302.000
IC
        [6]
        ICM: C07D239-54
        ICS: A61K031-52
        514/242; 514/243; 514/269; 544/298; 544/299
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 59 OF 78 USPATFULL on STN
        1998:104752 USPATFULL
ΑN
        Amine substituted compounds
TI
        Klein, J. Peter, Vashon, WA, United States
Underiner, Gail E., Brier, WA, United States
ΙN
        Kumar, Anil M., Seattle, WA, United States
        Ridgers, Lance H., Bothell, WA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
                                   19980901
ΡI
        US 5801182
        US 1995-485777
                                   19950607 (8)
ΑI
        Continuation-in-part of Ser. No. US 1994-217051, filed on 24 Mar 1994,
RLI
        now_abandoned
DT
        Utility
FS
        Granted
LN.CNT 1706
INCL
        INCLM: 514/269.000
        INCLS: 514/274.000; 544/310.000; 544/311.000; 544/312.000
NCL
                514/269.000
        NCLM:
        NCLS:
                514/274.000; 544/310.000; 544/311.000; 544/312.000
IC
        [6]
        ICM: A61K031-505
        ICS: C07D239-02
EXF
        544/312; 514/269; 514/274; 514/310; 514/311
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 60 OF 78 USPATFULL on STN
        1998:104751 USPATFULL
AN
        Amino alcohol substituted cyclic compounds
TI
        Michnick, John, Seattle, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
        Klein, J. Peter, Vashon Island, WA, United States
Rice, Glenn C., Seattle, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 5801181
PA
        us 5801181
                                   19980901
PΙ
        US 1995-474820
                                   19950607 (8)
ΑI
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Division of Ser. No. US 1993-152650, filed on 12 Nov 1993, now abandoned
RLI
        which is a continuation-in-part of Ser. No. US 1993-40820, filed on 31
        Mar 1993
        Utility
DT
        Granted
FS
LN.CNT 2822
        INCLM: 514/263.000
INCL
        INCLS: 514/183.000; 514/249.000; 514/259.000; 514/274.000; 514/309.000;
                514/315.000; 514/418.000; 514/425.000; 514/617.000; 514/619.000;
                514/626.000; 514/668.000; 514/669.000
                514/263.350
NCL
        NCLM:
                514/183.000; 514/249.000; 514/266.300; 514/274.000; 514/309.000; 514/315.000; 514/418.000; 514/425.000; 514/617.000; 514/619.000; 514/626.000; 514/668.000; 514/669.000
        NCLS:
IC
        [6]
        ICM: A01N043-00
        ICS: A01N043-90; A01N043-58; A01N043-42
        514/263; 514/249; 514/259; 514/265; 514/274; 514/309; 514/315; 514/418; 514/425; 514/617; 514/619; 514/626; 514/668; 514/669
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 61 OF 78 USPATFULL ON STN
L7
        1998:98932 USPATFULL
ΑN
        DHA-pharmaceutical agent conjugates of taxanes
TI
        Shashoua, Victor E., Brookline, MA, United States
IN
        Swindell, Charles S., Merion, PA, United States
        Webb, Nigel L., Bryn Mawr, PA, United States
        Bradley, Matthews O., Laytonsville, MD, United States
        Neuromedica, Inc., Conshohocken, PA, United States (U.S. corporation)
PA
        us 5795909
                                    19980818
PΙ
        us 1996-651312
ΑI
                                    19960522 (8)
        Utility
DT
        Granted
FS
LN.CNT 2451
INCL
        INCLM: 514/449.000
        INCLS: 514/549.000
                514/449.000
NCL
        NCLM:
                514/549.000
        NCLS:
IC
        [6]
        ICM: A61K031-335
        ICS: A61K031-22
        514/449; 514/549
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 62 OF 78 USPATFULL on STN
        1998:88470 USPATFULL
ΑN
          ***VEGF***
ΤI
                         gene transfer into endothelial cells for vascular
        prosthesis
        Pratt, Richard E., Palo Alto, CA, United States
Dzau, Victor J., Los Altos Hills, CA, United States
IN
PA
        The Board of Trustees of the Leland Stanford Junior Univ., Palo Alto,
        CA, United States (U.S. corporation)
US 5785965 19980728
PΙ
                                                                               <--
        US 1996-647821
                                    19960515 (8)
ΑI
        Utility
DT
        Granted
FS
LN.CNT 905
INCL
        INCLM: 424/093.210
        INCLS: 424/093.100; 424/093.200; 435/172.300; 435/325.000
                424/093.210
NCL
                424/093.100; 424/093.200; 435/325.000; 435/455.000; 435/456.000
        NCLS:
IC
        [6]
        ICM: A01N063-00
        ICS: C12N015-00
        600/36; 623/1; 623/11; 623/12; 435/172.3; 435/240.2; 435/320.1; 435/325; 424/93.21; 424/93.2; 514/44
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 63 OF 78 USPATFULL ON STN
L7
        1998:82763 USPATFULL
ΑN
TI
        Hydroxyl-containing xanthine compounds
        Underiner, Gail E., Brier, WA, United States
IN
        Porubek, David, Seattle, WA, United States
Klein, J. Peter, Vashon Island, WA, United States
        Woodson, Paul, Edmonds, WA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
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PA

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19980714
ΡI
        us 5780476
                                    19950606 (8)
        US 1995-468660
ΑI
RLI
        Division of Ser. No. US 1993-153256, filed on 16 Nov 1993, now abandoned
        which is a continuation-in-part of Ser. No. US 1992-976353, filed on 16
        Nov 1992, now patented, Pat. No. US 5473070
DT
        Utility
        Granted
FS
LN.CNT 1672
INCL
        INCLM: 514/263.000
        INCLS: 544/267.000
NCL
        NCLM:
               514/263.360
IC
        [6]
        ICM: A61K031-52
        ICS: C07D473-04
        514/263; 514/256; 514/257; 514/258; 514/259; 514/261; 514/269; 514/270
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 64 OF 78 USPATFULL on STN
        1998:79344 USPATFULL
ΑN
        Method for preparing substituted amino alcohol compounds
TI
        Klein, J. Peter, Vashon, WA, United States
TN
        Underiner, Gail E., Brier, WA, United States
        Kumar, Anil M., Seattle, WA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PA
                                    19980707
ΡI
        us 5777117
        us 1995-472569
                                    19950607 (8)
ΑI
        Division of Ser. No. US 1994-303842, filed on 8 Sep 1994 which is a continuation-in-part of Ser. No. US 1993-152650, filed on 12 Nov 1993 And Ser. No. US 1993-164081, filed on 8 Dec 1993 which is a continuation-in-part of Ser. No. US 1993-40820, filed on 31 Mar 1993,
RLI
        now abandoned, said Ser. No. US -152650 which is a
        continuation-in-part of Ser. No. US
                                                  -40820
DT
        Utility
        Granted
FS
LN.CNT 3153
INCL
        INCLM: 544/267.000
        INCLS: 544/257.000; 544/285.000; 544/286.000; 544/287.000; 544/311.000;
                546/141.000; 546/243.000; 546/246.000; 548/477.000; 548/546.000
                544/267.000
NCL
        NCLM:
                544/257.000; 544/285.000; 544/286.000; 544/287.000; 544/311.000; 546/141.000; 546/243.000; 546/246.000; 548/477.000; 548/546.000
        NCLS:
IC
        ICM: C07D473-10
        ICS: C07D239-80; C07D211-94; C07D209-48
        544/267; 544/257; 544/285; 544/286; 544/287; 544/311; 546/141; 546/243; 546/246; 548/477; 548/546
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 65 OF 78 USPATFULL on STN
ΑN
        1998:79342 USPATFULL
TI
        Acetal-and ketal-substituted pyrimidine compounds
        Leigh, Alistair, Brier, WA, United States
Underiner, Gail, Brier, WA, United States
ΙN
PA
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
ΡI
        US 5777115
                                    19980707
AΙ
        us 1994-193331
                                    19940207 (8)
RLI
        Continuation-in-part of Ser. No. US 1993-4353, filed on 14 Jan 1993, now
        abandoned
DT
        Utility
FS
        Granted
LN.CNT 1632
INCL
        INCLM: 544/242.000
        INCLS: 544/267.000; 514/269.000; 514/270.000; 514/256.000
NCL
                544/242.000
        NCLM:
                544/267.000
        NCLS:
IC
        [6]
        ICM: C07D239-26
        ICS: A61K031-505
        544/267; 544/242; 546/242; 546/243; 514/256; 514/269; 514/270
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 66 OF 78 USPATFULL on STN
L7
        1998:72634 USPATFULL
ΑN
        Prevention and treatment of cardiovascular pathologies
TI
IN
        Grainger, David J., Cambridge, England
        Metcalfe, James C., Cambridge, England
```

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Kunz, Lawrence L., Redmond, WA, United States
        Schroff, Robert W., Edmonds, WA, United States
       Weissberg, Peter L., Cambridge, England
NeoRx Corporation, Seattle, WA, United States (U.S. corporation)
PΑ
                                  19980623
PΙ
        US 5770609
       US 1995-486334
ΑI
                                  19950607 (8)
RLI
        Continuation-in-part of Ser. No. US 1994-242161, filed on 12 May 1994
       which is a continuation-in-part of Ser. No. US 1993-61714, filed on 13
       May 1993, now abandoned And a continuation-in-part of Ser. No. US
        1994-241844, filed on 12 May 1994 which is a continuation-in-part of
       Ser. No. US 1993-62451, filed on 13 May 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-11669, filed on 28 Jan 1993,
        now abandoned
DT
        Utility
        Granted
FS
LN.CNT 4318
INCL
        INCLM: 514/319.000
        INCLS: 514/324.000; 514/422.000; 514/428.000; 514/444.000; 514/448.000;
               514/651.000
NCL
       NCLM:
               514/319.000
       NCLS:
               514/324.000; 514/422.000; 514/428.000; 514/444.000; 514/448.000;
               514/651.000
IC
        [6]
        ICM: A61K031-445
        ICS: A61K031-40; A61K031-38; A61K031-135
514/651; 514/324; 514/212; 514/422; 514/428; 514/444; 514/448; 514/319
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 67 OF 78 USPATFULL on STN
ΑN
        1998:72620 USPATFULL
TI
        Oxime substituted therapeutic compounds
IN
       Klein, J. Peter, Vashon, WA, United States
        Leigh, Alistair, Brier, WA, United States
PΑ
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
                                  19980623
PΙ
       us 5770595
       US 1994-193344
                                  19940207 (8)
ΑI
RLI
       Continuation of Ser. No. US 1993-6083, filed on 19 Jan 1993, now
        abandoned
DT
       Utility
FS
       Granted
LN.CNT 2183
INCL
        INCLM: 514/263.000
        INCLS: 544/271.000; 544/273.000
NCL
       NCLM:
               514/263.350
       NCLS:
               514/151.000; 544/271.000; 544/273.000
IC
        [6]
       ICM: M61K031-52
EXF
        514/263; 544/271; 544/273
L7
     ANSWER 68 OF 78 USPATFULL ON STN
        1998:51651 USPATFULL
ΑN
       Substituted amino alcohol compounds
TT
IN
       Klein, J. Peter, Vashon, WA, United States
       Underiner,
                   Gail E., Brier, WA, United States
       Kumar, Anil M., Seattle, WA, United States
PA
       Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
PΙ
       us 5750575
                                  19980512
ΑI
       US 1995-475721
                                  19950607 (8)
       Division of Ser. No. US 1994-303842, filed on 8 Sep 1994, now patented,
RLI
       Pat. No. US 5641783 which is a continuation-in-part of Ser. No. US
        1993-152650, filed on 12 Nov 1993 And a continuation-in-part of Ser. No.
       US 1993-164081, filed on 8 Dec 1993, now patented, Pat. No. US 5470878
       which is a continuation-in-part of Ser. No. US 1993-40820, filed on 31
       Mar 1993, now abandoned
       Utility
DT
       Granted
LN.CNT 3115
INCL
       INCLM: 514/617.000
       INCLS: 514/653.000; 564/182.000; 564/355.000; 564/361.000
NCL
       NCLM:
               514/617.000
       NCLS:
               514/653.000; 564/182.000; 564/355.000; 564/361.000
IC
       [6]
       ICM: A61K031-165
       ICS: A61K031-135; C07C233-35; C07C215-20 564/355; 564/182; 564/361; 514/617; 514/653
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
```

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L7
      ANSWER 69 OF 78 USPATFULL on STN
         97:114932 USPATFULL
AN
         Suppression of nitric oxide production by osteopontin
TI
         Denhardt, David T., Bridgewater, NJ, United States
IN
         Hwang, Shiaw-Min, Piscataway, NJ, United States
Heck, Diane Elaine, Rumson, NJ, United States
         Lopez, Cecilia Ang, North Brunswick, NJ, United States
         Laskin, Debra L., Basking Ridge, NJ, United States
         Laskin, Jeffrey D., Piscataway, NJ, United States
         Rutgers University, Piscataway, NJ, United States (U.S. corporation)
PA
         University of Medicine & Dentistry of NJ, Newark, NJ, United States
         (U.S. corporation)
         us 5695761
PΙ
                                        19971209
                                                                                       <--
         US 1993-173116
                                        19931223 (8)
ΑI
         Utility
DT
FS
         Granted
LN.CNT 1552
         INCLM: 424/184.100
INCL
         INCLS: 424/085.500; 424/278.100; 530/351.000; 530/330.000; 530/326.000;
                  530/300.000; 514/002.000; 514/012.000
NCL
         NCLM:
                  424/184.100
                  424/085.500; 424/278.100; 514/002.000; 514/012.000; 530/300.000;
         NCLS:
                  530/326.000; 530/330.000; 530/351.000
IC
         [6]
         ICM: A01N037-18
ICS: A61K038-00; A61K039-38; C07K002-00

EXF 424/88; 424/85.5; 424/278.1; 424/184.1; 530/351; 530/330; 530/326; 530/300; 514/2; 514/12

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 70 OF 78 USPATFULL ON STN
         97:86614 USPATFULL
ΑN
         Halogen, isothiocyanate or azide substituted xanthines
TI
IN
         Leigh, Alistair, Brier, WA, United States
         Michnick, John, Seattle, WA, United States
         Kumar, Anil, Seattle, WA, United States
         Underiner, Gail, Brier, WA, United States
         Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation) US 5670506 19970923 <--
PA
ΡI
         us 1993-42946
ΑI
                                        19930405 (8)
         Utility
DT
FS
         Granted
LN.CNT 1994
INCL
         INCLM: 514/258.000
         INCLS: 514/263.000; 544/267.000; 544/272.000; 544/277.000
NCL
                  514/141.000
                  544/267.000; 544/272.000; 544/277.000
         NCLS:
IC
         [6]
         ICM: A61K031-52
         ICS: C07D473-00
EXF 544/267; 544/276; 544/272; 544/277; 514/258 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 71 OF 78 USPATFULL on STN
L7
         97:73601 USPATFULL
AN
TI
         Compositions for inhibiting restenosis
         Weisz, Paul B., State College, PA, United States
The Trustees of the University of Pennsylvania, Philadephia, PA, United
IN
PA
         States (U.S. corporation)
                                        19970819
         us 5658894
PΙ
         us 1994-345011
                                        19941123 (8)
ΑI
         Continuation of Ser. No. US 1992-900592, filed on 18 Jun 1992, now abandoned And a continuation-in-part of Ser. No. US 1991-790320, filed on 12 Nov 1991, now abandoned which is a continuation-in-part of Ser. No. US 1991-691168, filed on 24 Apr 1991, now abandoned which is a continuation of Ser. No. US 1989-397559, filed on 23 Aug 1989, now
RLI
         abandoned , said Ser. No. US
                                              -900592 which is a continuation-in-part
         of Ser. No. US 1990-480407, filed on 15 Feb 1990, now patented, Pat. No.
         US 5183809, issued on 2 Feb 1993
         Utility
DT
FS
         Granted
LN.CNT 1449
         INCLM: 514/058.000
INCL
         INCLS: 514/021.000; 514/023.000; 514/054.000; 514/060.000; 536/103.000; 530/810.000; 530/812.000; 530/813.000
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NCL
        NCLM:
                514/058.000
                514/021.000; 514/023.000; 514/054.000; 514/060.000; 530/810.000; 530/813.000; 536/103.000
        NCLS:
IC
        [6]
        ICM: A61K031-735
        ICS: C08B037-16
        514/21; 514/23; 514/54; 514/58; 514/60; 536/103; 530/810; 530/812;
EXF
        530/813
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 72 OF 78 USPATFULL ON STN
        97:54233 USPATFULL
ΑN
TI
        Substituted amino alcohol compounds
        Klein, J. Peter, Vashon, WA, United States
Underiner, Gail E., Brier, WA, United States
IN
        Kumar, Anil M., Seattle, WA, United States
        Cell Therapeutics, Inc., Seattle, WA, United States (U.S. corporation)
US 5641783 19970624 <--
PA
PI
                                    19940908 (8)
ΑI
        us 1994-303842
RLI
        Continuation-in-part of Ser. No. US 1993-152650, filed on 12 Nov 1993
        And Ser. No. US 1993-164081, filed on 8 Dec 1993, now patented, Pat. No.
        us 5470878
DT
        Utility
FS
        Granted
        3206
LN.CNT
INCL
        INCLM:
                514/263.000
                514/183.000;
                               514/222.500; 514/223.500;
514/228.800; 514/229.200;
        INCLS:
                                                             514/224.200;
514/230.500;
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                514/227.500;
                                                                            514/230.800
                514/237.800;
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                514/247.000;
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                                                             514/255.000;
                                                                            514/256.000
                514/258.000; 514/259.000;
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                                                             514/262.000;
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                               514/274.000;
                                              514/297.000;
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                                              514/306.000; 514/307.000;
                                                                            514/311.000:
                514/302.000;
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                                                                            514/357.000;
                514/359.000; 514/360.000; 514/361.000;
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                                                             514/369.000;
                514/364.000;
                              514/365.000; 514/367.000;
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                514/373.000;
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514/399.000;
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514/395.000;
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514/398.000;
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                               514/406.000;
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                               514/423.000;
                                                                            514/427.000
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                514/428.000:
                                              544/002.000;
                                                             544/003.000;
                544/053.000;
                               544/063.000;
                                              544/065.000;
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                                                             544/066.000;
                               544/091.000;
                                              544/162.000;
                                                             544/215.000;
                544/090.000;
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                               544/220.000;
                                              544/224.000;
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                544/219.000;
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                544/254.000;
                               544/255.000;
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                                                             544/262.000;
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                544/277.000;
                               544/278.000;
                                              544/280.000;
                                                             544/283.000; 544/286.000;
                               544/311.000;
                544/301.000;
                                              544/335.000;
                                                             544/336.000; 544/350.000;
                               544/385.000;
                544/353.000;
                                              544/401.000;
                                                             546/102.000;
                                                                            546/113.000
                               546/115.000;
                                              546/117.000;
546/139.000;
546/176.000;
                546/114.000;
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                546/122.000;
546/157.000;
                               546/138.000;
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546/178.000;
                                                                            546/153.000
                               546/164.000;
                                                                            546/242.000
                546/243.000;
                               546/246.000;
                                              546/264.000;
                                                             546/300.000;
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548/179.000;
                               548/123.000:
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                548/100.000;
                               548/134.000;
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                548/221.000;
                               548/225.000;
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                548/237.000;
                               548/240.000;
                                              548/241.000;
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                548/252.000;
                                              548/267.800;
                                                             548/303.700;
                               548/267.200;
                                                                            548/306.400;
                               548/309.700;
                548/307.100;
                                              548/319.100;
                                                             548/323.500;
                                                                            548/340.100;
                                              548/356.100;
                                                             548/370.100;
                548/348.100;
                               548/349.100;
                                                                            548/375.100;
                               548/452.000;
548/486.000;
                548/379.400;
                                              548/453.000;
                                                             548/470.000;
                                                                            548/482.000;
                548/485.000;
                                              548/491.000;
                                                             548/503.000;
                                                                            548/532.000;
                548/543.000;
                               548/546.000;
                                              548/550.000; 548/565.000; 548/566.000
                514/263.350
NCL
        NCLM:
        NCLS:
                514/183.000;
                               514/222.500;
                                              514/223.500;
                                                             514/224.200; 514/226.800;
                514/227.500:
                               514/228.800:
                                              514/229.200;
                                                             514/230.500;
                                                                            514/230.800
                514/237.800;
                               514/241.000;
                                              514/242.000;
                                                             514/243.000;
                                                                            514/246.000
                514/247.000;
                               514/248.000;
                                              514/249.000;
                                                                            514/256.000
                                                             514/252.160;
                               514/264.100;
                                              514/266.300;
                514/259.500;
                                                                            514/274.000;
                                                             514/270.000:
                514/297.000;
                               514/300.000;
                                              514/301.000;
                                                             514/302.000;
                                                                            514/303.000;
                               514/307.000;
                514/306.000
                                              514/311.000;
                                                             514/312.000;
                                                                            514/315.000;
                514/345.000
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                548/267.800;
                               548/323.500; 548/340.100; 548/348.100; 548/349.100;
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548/550.000; 548/565.000; 548/566.000
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        ICM: A61K031-415
        ICS: A61K031-42; A61K031-425; A61K031-52
        544/272; 514/263
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 73 OF 78 USPATFULL on STN
ΑN
        97:5708 USPATFULL
        Method for identifying an agent which increases TGF-beta levels
TI
        Grainger, David J., Cambridge, England
IN
        Metcalfe, James C., Cambridge, England
NeoRx Corporation, Seattle, WA, United States (U.S. corporation)
US 5595722 19970121 <--
PA
PΙ
        us 1995-476735
ΑI
                                    19950607 (8)
        Continuation-in-part of Ser. No. US 1994-242161, filed on 12 May 1994
RLI
        which is a continuation-in-part of Ser. No. US 1993-61714, filed on 13
        May 1993, now abandoned And Ser. No. US 1994-241844, filed on 12 May
        1994 which is a continuation-in-part of Ser. No. US 1993-62451, filed on
        13 May 1993, now abandoned which is a continuation-in-part of Ser. No.
        US 1993-11669, filed on 28 Jan 1993, now abandoned
DT
        Utility
FS
        Granted
LN.CNT 4090
INCL
        INCLM: 424/009.200
        NCLM: 424/009.200
NCL
IC
        [6]
        ICM: A61K049-00
        424/9.2
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 74 OF 78 USPATFULL ON STN
AN
        96:53051 USPATFULL
        Extraluminal regulation of the growth and repair of tubular structures
TI
IN
        Edelman, Elazer R., Brookline, MA, United States
        Adams, David H., Boston, MA, United States
        Karnovsky, Morris J., Newton Centre, MA, United States
President and Fellows of Harvard College, Cambridge, MA, United States
PA
        (U.S. corporation)
        us 5527532
PΙ
                                    19960618
                                    19930902 (8)
        us 1993-105903
ΑI
        Continuation-in-part of Ser. No. US 1991-656182, filed on 27 Feb 1991,
RLI
        now abandoned which is a continuation-in-part of Ser. No. US
        1989-436337, filed on 13 Nov 1989, now abandoned
        Utility
DT
        Granted
FS
LN.CNT 940
        INCLM: 424/422.000
INCL
        INCLS: 424/423.000; 424/426.000; 424/430.000
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NCL
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                  424/422.000
         NCLS:
                  424/423.000; 424/426.000; 424/430.000
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IC
         ICM: A61K009-12
         424/422; 424/423; 424/426; 424/430; 514/56; 514/423; 514/12
EXF
L7
      ANSWER 75 OF 78 USPATFULL ON STN
         95:105868 USPATFULL
ΑN
         Cell signaling inhibitors
ΤI
        Michnick, John, Seattle, WA, United States
Underiner, Gail E., Brier, WA, United States
Klein, J. Peter, Vashon Island, WA, United States
Rice, Glenn C., Seattle, WA, United States
Cell Therapeutics, Inc., Seattle, WA, United States
US 5470878 19951128 <--
IN
PA
ΡI
         US 1993-164081
                                         19931208 (8)
ΑI
RLI
         Continuation-in-part of Ser. No. US 1993-40820, filed on 31 Mar 1993,
         now abandoned
DT
         Utility
FS
         Granted
LN.CNT 2665
         INCLM: 514/558.000
INCL
         INCLS: 514/258.000; 514/262.000; 514/274.000; 514/299.000; 514/315.000; 514/418.000; 514/425.000; 514/529.000; 514/552.000; 514/561.000; 514/613.000; 514/617.000; 514/626.000; 514/629.000; 514/669.000; 544/254.000; 544/285.000; 544/301.000; 546/183.000; 546/243.000;
                                    548/556.000; 554/055.000; 554/061.000;
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                   554/213.000;
                                    560/130.000; 560/145.000; 562/553.000;
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         NCLS:
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                   544/301.000; 546/183.000; 546/243.000; 548/486.000; 548/556.000
IC
         [6]
         ICM: A61K031-20
         ICS: C07C233-00

554/51; 554/61; 554/55; 554/108; 554/213; 564/224; 564/506; 564/198;

564/215; 564/201; 564/197; 514/625; 514/629; 514/613; 514/558; 514/552;

514/529; 514/561; 514/626; 514/669; 560/130; 560/145; 562/553; 562/567
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 76 OF 78 USPATFULL ON STN
ΑN
         94:93338 USPATFULL
TI
         Methods for treating arteriosclerosis
TN
         Halperin, Jose, Brookline, MA, United States
         Brugnara, Carlo, Newton Highlands, MA, United States
PΑ
         President and Fellows of Harvard University, Cambridge, MA, United
         States (U.S. corporation)
         US 5358959
US 1993-18835
ΡI
                                         19941025
                                                                                           <--
ΑI
                                         19930218 (8)
         Utility
DT
FS
         Granted
LN.CNT 539
INCL
         INCLM: 514/396.000
         INCLS: 514/399.000; 514/824.000
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         NCLS:
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IC
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         ICM: A61K031-415
EXF
         514/396; 514/399; 514/824
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 77 OF 78 USPATFULL ON STN
L7
AN
         91:84437 USPATFULL
TI
         Method for preventing tissue damage after an ischemic episode
IN
         Sheffield, Warren D., Lebanon, NJ, United States
         Ethicon, Inc., Somerville, NJ, United States (U.S. corporation)
PA
PI
         us 5057494
                                         19911015
ΑI
         US 1988-227579
                                         19880803 (7)
         Utility
DT
FS
         Granted
LN.CNT 487
         INCLM: 514/012.000
INCL
         INCLS: 514/021.000
         NCLM: 514/012.000
NCL
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NCLS: 514/021.000
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       ICM: A61K037-02
       ICS: A61K037-36
       514/12; 514/21
FXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 78 OF 78 WPIDS COPYRIGHT 2004 THE THOMSON CORP on STN
     2000-256866 [22]
ΑN
                         WPIDS
     C2000-078440
DNC
     Hydrogel compositions useful for controlled delivery of growth factors
TI
     e.g. in treatment of ischemia and in wound healing.
DC
     A11 A25 A96 B04 B07
     JENNINGS, R N; PROTTER, A A; WANG, Y J; YANG, B
IN
     (SCIO-N) SCIOS INC
PA
     87
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     wo 2000013710
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             GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU
             LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR
             TT UA UG US UZ VN YU ZA ZW
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A2 20010620 (200135)
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     EP 1107791
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                                                          A61K047-10
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     AU 758178
                      в 20030320 (200329)
                                                           A61K047-10
     WO 2000013710 A2 WO 1999-US20382 19990903; AU 9959095 A AU 1999-59095
ADT
     19990903; EP 1107791 A2 EP 1999-946759 19990903, WO 1999-US20382 19990903;
     US 6331309 B1 Provisional US 1998-99168P 19980904, US 1999-390164
     19990903; JP 2002524425 w wo 1999-US20382 19990903, JP 2000-568516
     19990903; AU 758178 B AU 1999-59095 19990903
FDT
     AU 9959095 A Based on WO 2000013710; EP 1107791 A2 Based on WO 2000013710;
     JP 2002524425 W Based on WO 2000013710; AU 758178 B Previous Publ. AU
9959095, Based on WO 2000013710
PRAI US 1998-99168P 19980904;
                            19980904; US 1999-390164
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          A61F013-00; A61K038-22; A61K047-10
A61K009-10; A61K009-70; A61K038-18; A61K047-26; A61K047-32;
A61K047-34; A61K047-36; A61P009-10; A61P017-02
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